

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
19 September 2002 (19.09.2002)

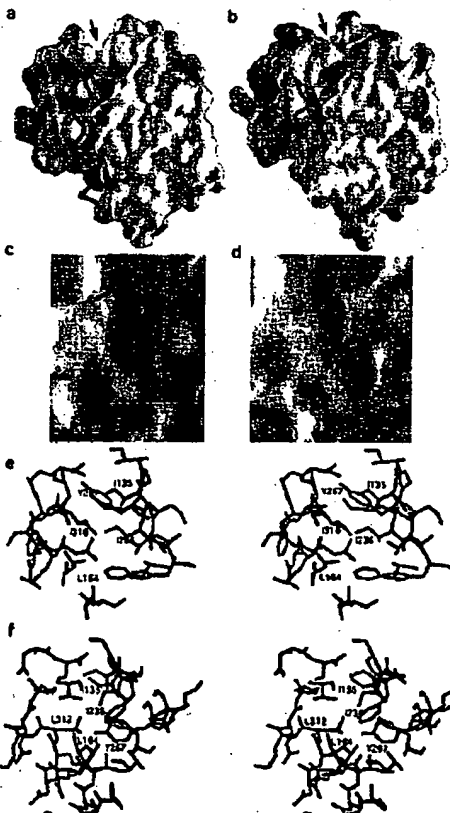
PCT

(10) International Publication Number
WO 02/072030 A2

- (51) International Patent Classification⁷: A61K (72) Inventors; and
(21) International Application Number: PCT/US02/07809 (75) Inventors/Applicants (for US only): ARNAOUT, M., Amin [US/US]; 99 Shaw Road, Chestnut Hill, MA 02467 (US). LI, Rui [CN/US]; 89th Street, Apartment 107, Medford, MA 02155 (US). XIONG, Jian-Ping [CN/US]; 95 W. Squantum Street, #1105, Quincy, MA 02171 (US).
(22) International Filing Date: 13 March 2002 (13.03.2002)
(25) Filing Language: English
(26) Publication Language: English (74) Agent: MEIKLEJOHN, Anita, L.; Fish & Richardson P.C., 225 Franklin Street, Boston, MA 02110 (US).
(30) Priority Data: 09/805,354 13 March 2001 (13.03.2001) US (81) Designated States (national): AE, AG, AI, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.
(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application: US 09/805,354 (CIP) Filed on 13 March 2001 (13.03.2001)
(71) Applicant (for all designated States except US): THE GENERAL HOSPITAL CORPORATION [US/US]; 55 Fruit Street, Boston, MA 02114 (US).

[Continued on next page]

(54) Title: HIGH AFFINITY INTEGRIN POLYPEPTIDES AND USES THEREOF



(57) Abstract: Polypeptides comprising all or part of a variant integrin α subunit A domain or a variant integrin β subunit A-like domain are described. In solution or in membrane-associated form, the A domain or the A-like domain of the polypeptides of the invention exists predominantly in a high affinity conformation. In the polypeptides of the invention, referred to as variant integrin polypeptides, a crucial isoleucine residue (described in greater detail below) is absent. The isoleucine can be either deleted or replaced with different amino acids residue, preferably a smaller or less hydrophobic amino acid residue, e.g., alanine or glycine. Because the variant integrin polypeptides of the invention exist in solution or in membrane-associated form predominantly in a high affinity conformation, they are useful in screening assays for the identification of molecules that bind to (and/or mediate the activity of) an integrin. They are also useful for generating antibodies, e.g., monoclonal antibodies, which bind to the high affinity form of an integrin. Some such antibodies recognize an epitope that is either not present or not accessible on an integrin that is in a lower affinity conformation. The variant integrin polypeptides of the invention can be derived from any integrin α subunit or any integrin β subunit and could be used therapeutically. The variant integrin polypeptides preferably include a ligand-binding portion of an A-domain or an A-like domain.

WO 02/072030 A2

BEST AVAILABLE COPY



(84) Designated States (*regional*): ARIPO patent (GH, GM, KI, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

5 **HIGH AFFINITY INTEGRIN POLYPEPTIDES AND USES**
 THEREOF

10

BACKGROUND

 Integrins are heterodimeric receptors that mediate a wide variety of important interactions both between cells and between cells and the extracellular matrix via ligand
15 binding. All integrins have an α subunit and a β subunit. Within the α subunit a region referred to as the A domain (or I domain) is known to be an important mediator of ligand binding. A similar region, the A-like domain, is present in many β subunits. Many integrins are thought to exist in two conformations, a low affinity state (the "closed" or "unliganded" conformation") and a high affinity state (the "open" or "liganded"
20 conformation), the later of which is responsible for high affinity ligand binding.

 Integrins transduce signals that mediate the effects of the matrix on the physiological activity of cells (e.g., motility, proliferation, differentiation). Moreover, integrins play a role in inflammation and in oncogenic cell transformation, metastasis, and apoptosis. Thus, there is considerable interest in identifying compounds that can
25 activate or inhibit the activity of one or more integrins.

 In order for an efficient integrin-ligand binding to occur, it is thought that the integrin must be in its high affinity configuration. It appears that inside-out signals generated when cells are activated by a variety of stimuli apparently switch integrins from a low affinity state to a high affinity state. This functional upregulation is
30 associated with conformational changes in the extracellular regions of integrins that

- 5 include the A domain of the α -subunit and the A-like domain of the β subunit (Smith et al. 1988 *J. Biol. Chem.* 263:18726).

The integrin A-domain assumes a dinucleotide-binding fold (Lee et al. 1995 *Cell* 80:631; Emsley et al. 1997 *J. Biol. Chem.* 272:28512; Li et al. 1998 *J. Cell Biol.* 143:1523; Nolte et al. 1999 *FEBS Lett.* 452:379; and Legge et al. 2000 *J. Mol. Biol.* 10 295:1251), with a metal ion dependent adhesion site (MIDAS) on its top, and is connected through a C-terminal α 7 helix at its bottom to the body of the integrin. MIDAS and its surrounding exposed side-chains form the binding site for physiologic ligands (Li et al., *supra*; Michishita et al., *Cell* 72:857-867, 1993; Kamata et al., *J. Biol. Chem.* 269:26006-26010, 1994; Kern et al., *J. Biol. Chem.* 269:22811-6, 1994; Edwards 15 et al., *J Biol. Chem.* 273:28937-44, 1998; Zhang et al., *Biochemistry* 38:8064-71, 1999) and certain antagonists (Rieu et al., *J Biol Chem.* 271:15858-15861, 1996). In the "open" conformation, three non-charged residues in the protein directly coordinate the metal ion in MIDAS, a pseudoligand or ligand glutamate residue (Lee et al., *supra*; Li et al., *supra*; Emsley et al., *Cell* 100:47-56, 2000) completes metal coordination. In the "closed" form, 20 the amphipathic C-terminal α 7 helix is shifted upwards by 10Å compared to the "open" form, wrapping around the rest of the domain. This large shift is associated with a change in metal coordination, where one of the three coordinating residues, a threonine, is now replaced with an aspartate, and a water molecule replaces the glutamate in completing the metal ion coordination sphere (Lee et al., *Structure* 3:1333-1340, 1995). 25 These changes in metal coordination and topology of MIDAS are similar to those described in the structurally homologous G proteins (Lee et al., *supra*).

The crystal structure of four integrin A-domains (CD11b, CD11a, CD49a and CD49b) have been reported to date (Lee et al., *supra*; Lee et al., *Structure* 3:1333, 1995; Emsley et al., *supra*; Li et al., *supra*; Emsley et al., *J. Biol. Chem.* 272:28512, 1997). All 30 with the exception of integrin CD11b A-domain (11bA), were found only in the "closed" form, leading to the suggestion that the "open" form is a non-informative crystal artifact (Baldwin et al., *Structure* 6:923-935, 1998). Three studies support the view that the "open" form of the integrin A-domain equates with the "high" affinity state (Li et al.

5 1998 *J. Cell Biol* 143:1523; Rieu et al., *supra*; Oxvig et al., *Proc. Nat'l. Acad. Sci. USA* 96:2215-20, 1999). In the first, point mutations in CD11bA that are predicted on structural grounds to destabilize the "closed" structure, increased the proportion of the "high affinity" form in solution (Li et al. 1998 *J. Cell Biol.* 143:1523). In the second, the binding site for an "activation-dependent" monoclonal antibody mapped to a
10 conformationally sensitive region of the A-domain (Oxvig et al., *supra*). The third study showed that an A-domain in complex with a short collagen peptide assumed the "open" conformation, and suggested that the "open" form can only be obtained in the presence of ligand (Emsley et al., *supra*). While it has been suggested that the ligand causes the conformational change in integrins, at least one study suggests that integrins can exist in
15 high affinity state even in the absence of ligand (Smith and Cheresch 1988 *J Biol Chem* 263:18726-31). In addition, several studies suggested that ligand binding affinity in heterodimeric integrins can be altered in an allosteric manner (Li et al. 1998 *J Cell Biol* 143:1523; Edwards et al, *J Biol Chem* 273:28937-28944, 1998; Calderwood et al, *J Biol Chem* 273:5625, 1998; Zhang et al., *J Biol Chem.* 271: 29953-7, 1996).

20

SUMMARY

The invention features polypeptides comprising all or part of a variant integrin α subunit A domain or a variant integrin β subunit A-like domain. In the polypeptides of the invention, referred to as variant integrin polypeptides, an important isoleucine residue (described in greater detail below) is absent. The isoleucine can be either deleted or
25 replaced with different amino acids residue, preferably a smaller or less hydrophobic amino acid residue, e.g., alanine or glycine. Because the variant integrin polypeptides of the invention tend to exist in solution in a high affinity conformation, they are useful in screening assays for the identification of molecules that bind to (and/or modulate the activity of) an integrin. They are also useful for generating antibodies, e.g., monoclonal
30 antibodies, which bind to the high affinity form of an integrin. Some such antibodies recognize an epitope that is either not present or not accessible on an integrin that is in a lower affinity conformation. Thus, the invention features antibodies which bind with

5 greater affinity to a variant integrin of the invention than to corresponding wild-type integrin. The variant integrin polypeptides of the invention can be derived from any integrin α subunit or any integrin β subunit. The variant integrin polypeptides preferably include a ligand-binding portion of an A-domain or an A-like domain.

The invention also features methods for identifying a compound that binds to a
10 variant integrin polypeptide of the invention. Such screening methods can entail exposing the polypeptide to a test compound of interest and determining whether the compound binds to the polypeptide. Thus, the assay can be a simple binding assay (e.g., where binding of the compound is measured only in the absence of the ligand) or a competitive binding assay (e.g., where binding of the compound is measured in the
15 presence of the ligand). The invention also features methods for identifying a compound that interferes with the binding of an integrin ligand to an integrin by measuring the binding of an integrin ligand to a variant integrin polypeptide in the presence and absence of a test compound. The ability of a test compound to interfere with the binding of an integrin ligand to an integrin may also be tested. In addition, the binding specificity of a
20 compound can be assessed by measuring the binding of the integrin ligand to a second integrin (or measuring the binding of a second ligand to the integrin) in the presence and absence of the test compound.

The invention also features methods for interfering with the binding of an integrin to an integrin ligand by administering a variant integrin polypeptide of the invention or an
25 antibody that selectively binds to a variant integrin polypeptide of the invention.

The invention also features methods of administering a variant integrin polypeptide of the invention or an antibody that selectively binds to a variant integrin polypeptide of the invention for the purpose of identifying the presence of a ligand which binds to an active integrin. Such assays are useful for diagnosing inflammation, e.g.,
30 occult inflammation (e.g. abscess or an active atherosclerotic lesion).

The invention further features nucleic acid molecules (e.g., mRNA and DNA) encoding a polypeptide of the invention or a polypeptide which includes a polypeptide of the invention. The invention also includes nucleic acid molecule encoding a fusion

- 5 polypeptide comprising a polypeptide of the invention and a second polypeptide, e.g., an immunoglobulin constant domain.

The experiments described below concern the design a preparation of a variant form of CD11b (an integrin α subunit) that is more active than the wild-type form of CD11b. Without being bound by any particular theory, it appears that, in solution, the
10 amount of this variant subunit that is in the open (active) conformation is greater than for the corresponding wild-type form of the subunit.

Creation of the variant CD11b involves deletion or substitution of an invariable C-terminal Ile residue. Deletion or substitution of the Ile residue confers a high affinity phenotype in isolated CD11b A domain as well as in the intact integrin. Moreover, The
15 Ile-modified A-domain can be crystallized in the "open" conformation. Thus, without being bound by any particular theory, it appears that an Ile-based allosteric switch controls affinity and conformation in the integrin A-domain. Accordingly, variant, high affinity forms of other integrin α subunits (and β subunits) can be created by deleting or substituting the corresponding highly conserved Ile residue.

- 20 Stable open (high affinity) and closed (low affinity) forms of CD11b can also be created by introducing cysteine residues into CD11b so as to create a disulfide bridge that stabilizes the desired form of CD11b (or CD11b A domain). Thus, the invention also features a variant CD11b A domain-containing polypeptide in which F313 and A320 are replaced by cysteine residues (Amino acid positions refer to full-length CD11b starting
25 with a M, not mature CD11b, starting with an F.). This variant forms a stable open form of the A domain. The invention also features a variant CD11b A domain-containing polypeptide in which V315 and A320 are replaced by cysteine residues (Amino acid positions refer to full-length CD11b starting with a M, not mature CD11b, starting with an F.). This variant forms a stable closed form of the A domain. The invention also
30 features nucleic acid molecules encoding these variant forms, antibodies which bind selectively bind one or the other of these variant forms, antibodies which bind both variant forms, and methods for identifying compounds which selectively bind to one or

5 both of these variant forms. The invention also features A domain-containing peptides derived from CD11a, CD11c, CD11d, and other integrin subunits.

The details of one or more embodiments of the invention are set forth in the accompanying drawings and the description below. Other features, objects, and advantages of the invention will be apparent from the description and drawings, and from
10 the claims. All of the patents and publications referenced are herein incorporated by reference.

DESCRIPTION OF DRAWINGS

Figures 1A and 1B depict surface representation of the CD11bA domain crystal structure in its "closed" (Figure 1A) and "open" (Figure 1B) states, with the C-terminal
15 $\alpha 7$ helix outlined as a gray ribbon, the isoleucine (I316) residue in black. An arrow points to the MIDAS face. MIDAS and SILEN (socket for isoleucine) lie on almost opposite ends of the A-domain structure.

Figures 1C and 1D depict magnified face views of SILEN residues (I135, L164, I236, Y267, lying within 4Å radius from I316) in the "closed" (Figure 1C) and "open"
20 (Figure 1D) conformations. In the "closed" form (Figure 1C), I316 sidechain is wedged in SILEN; L312 is seen on top. L312 moves to cover SILEN in the "open" structure (Figure 1D). Figures 1A-1D were built using GRASP (Barry Honig, Columbia University, NY).

Figures 1E and 1F depict stereo views of the I316 coordination socket in the
25 "closed" (Figure 1E) and "open" (Figure 1F) conformations. SILEN residues and I316 and L312 are labeled.

Figures 2A and 2B depict $2F_0-F_C$ electron density maps of the C-terminal portions of $\alpha 7$ from 11bA¹²³⁻³²¹ (Figure 2A) and 11bA¹²³⁻³¹⁵ structures. The maps were contoured at 1.1 σ , and made with O.

Figures 2C and 2D depict ribbon diagrams showing the complete superimposition
30 of the 11bA¹²³⁻³²¹ (Figure 2C, red tracing) and 11bA¹²³⁻³¹⁵ (Figure 2D, red tracing) structures on the reported "closed" (Figure 2C, green tracing) and "open" (Figure 2D,

5 green tracing) forms of the CD11b A domain, respectively. The metal ion is shown as a purple sphere.

Figures 2E and 2F depict stereo views of MIDAS in the 11bA¹²³⁻³²¹ (Figure 2E) and 11bA¹²³⁻³¹⁵ (Figure 2F) structures. The direct D242-metal bond, and indirect T209-water-metal bond (Figure 2E) are characteristic of the "closed" conformation. The metal ion in this case is Mn²⁺. The MIDAS conformation (a direct T209-metal bond, indirect D242-water-metal bond, and a pseudo ligand glutamate, occupying the active site, shown in green, and directly coordinating the metal ion (Ca²⁺) are those of the "open" conformation). In Figures 2E and 2g, the metal ion is shown in purple, and hydrogen bonds as dotted yellow lines.

15 Figures 3A-3J depict the results of functional analysis of 11bA^{E-G} and 11bA¹²³⁻³¹⁵ A-domains using BIAcore™ for recording the interactions of 11bA^{E-G} (Figures 3A-3E) and 11bA¹²³⁻³¹⁵ (Figures 3F-3J) with the activation-dependent ligands iC3b (Figures 3A and 3F), fibrinogen (Figures 3B and 3G) or CD54 (Figures 3C and 3H). All three ligands bound to 11bA¹²³⁻³¹⁵, but not to 11bA¹²³⁻³²¹. The observed differences in binding were not
20 due to differences in protein concentration, as binding to the activation-independent ligands, NIF (Figures 3D and 3I) and mAb 904 (Figures 3E and 3J) were comparable. To quantitatively determine the affinity, various concentrations of the A domain peptides were used. The binding data were analyzed by linear transformation, giving dissociation constants (Kd, mean±SD, n=2) of 0.46±0.15 μM (for iC3b), 0.25±0.07 μM (for
25 fibrinogen); and 0.22±0.04 μM (for CD54).

Figure 4A depicts the results of an analysis of the binding of 11bA^{1→G} to iC3b. The dotted line represents the lack of binding of 11bA¹²³⁻³²¹ to the same ligand. The calculated Kd is 0.66±0.3 μM (mean ± SD, n=2).

Figure 4B depicts a histogram (mean ± SD, n=3) showing the relative binding of
30 1→G CR3 expressed on COS cells to iC3b, compared with wild-type receptor.

Figure 5 depicts an alignment of the A domains of nine alpha integrin α subunit (CD11b, CD11c, CD11d, CD11a, alpha 1, alpha 2, alpha 10, alpha 11, and alpha E). In this alignment, the invariant Ile (I316) is indicated by an arrow.

5 Figure 6 shows that an I to G mutation of the invariant isoleucine in CD11a A domain induces an activated domain in this integrin, underscoring the applicability of the findings in the CD11b A domain to other integrin A-domains.

 Figure 7 is an alignment of the A-like domains of eight integrin β subunits. In this alignment, the residue corresponding to the invariant Ile in β subunits is indicated by
 10 an arrow.

DETAILED DESCRIPTION

Example 1: Generation of a stable, high affinity CD11b variant A-domain by deletion

 A variant CD11b A domain consisting of amino acids E¹²³ to K³¹⁵ of CD11b (11bA¹²³⁻³¹⁵; also referred to as 11bA^{E-K}) was created. Note that the amino acid
 15 numbering in these example corresponds to the mature protein. While the amino acid numbering in Tables 2 and 3 below refers to the numbering in the complete protein (including signal sequence). CD11b has a 16 amino acid signal sequence. Thus, E¹²³ in the mature protein corresponds to E139 in the complete protein. This variant A domain was characterized and compared to a CD11b A domain consisting of amino acids E¹²³ to
 20 G³²¹ (11bA¹²³⁻³²¹; also referred to as 11bA^{E-G}). A variant A domain with a Ile to Gly change at residue 316 (11bA^{I→G}) was also created. All three proteins were expressed as GST fusion proteins that were cleaved to release the protein of interest.

 The variant polypeptides were created using standard recombinant techniques. Restriction and modification enzymes were purchased from New England Biolabs, Inc.
 25 (Beverly, MA), Boehringer Mannheim (Germany), or GIBCO BRL (Gaithersburg, MD). Site-directed mutagenesis was carried out in pGEX-4T-1 vector as described (Rieu et al. 1996 *J Biol Chem* 271:15858). The following mutagenic primers were used. IFAdel
 Fwd: 5'-TATAGGATCCGAGGCCCTCCGAGGGAGTCCTCAAGAGGATAG-3';
 Reverse: 5'-CTACTCGAGTTACTTCTCCCGAAGCTGGTTCTGAATGGTC-3'; I-G
 30 reverse: 5'-CTACTCGAGTTAACCCTCGATCGCAAAGCCCTTCTC-3'. Introduction of the respective mutation was confirmed by direct DNA sequencing. The PvuI-BspEI-restricted cDNA fragment of the A-domain containing the mutation was subcloned into

5 the PvuI-BspEI-restricted CD11b cDNA, cloned into pcDNA3 plasmid, which containing full-length human CD11b (Rieu et al. 1996 *J Biol Chem* 271:15858). 11b A¹²³⁻³²¹ and 11bA¹²³⁻³¹⁵ and 11bA^{1→G} A-domains were expressed as GST fusion proteins in *Escherichia coli* (Michishita et al. 1993 *Cell* 72:857), cleaved with thrombin and purified as described Li et al. 1999 *J. Cell Biol* 143:1523. C¹²⁹ was replaced by S in all the
10 expressed GST-A-domain fusion form to prevent formation of disulfide-linked dimmers in solution after thrombin cleavage (not shown). Purity was confirmed by SDS-PAGE analysis.

The structures of 11bA¹²³⁻³²¹ and 11bA¹²³⁻³¹⁵ were determined by x-ray crystallography. Crystals were grown using 10mg/ml stock protein solutions and the
15 hanging drop vapor diffusion method as described Li et al. (1998 *J. Cell Biol* 143:1523). Several crystal conditions were tried for each A-domain. 11bA¹²³⁻³¹⁵ and 11bA^{1→G} formed crystals in the presence of a reservoir solution containing 15% polyethylene glycol 8 K, 0.10 M Tris, pH 8.2, 150 mM CaCl₂, at room temperature. Crystals started to form within a week, grew to a typical size of 0.3 mm x 0.05 mm x 0.04 mm in two weeks, and
20 belonged to the tetragonal space group P4, with unit cell of a=b=45.7Å. 11bA¹²³⁻³²¹ did not crystallize under these conditions, but formed crystals at room temperature using 10% polyethylene glycol 4000, 0.1M sodium citrate, pH 4.5, 5 mM MnCl₂ in the reservoir buffer. 11bA¹²³⁻³²¹ was crystallized in the P₂₁2₁2₁ space group with unit cell of a=48.1 Å, b = 121.5 Å, c = 74.5 Å.

25 A single 11bA¹²³⁻³¹⁵ crystal was used to collect a 2.3 Å resolution data set, at 100 K, on beamline X12B of the National Synchrotron Light Source at the Brookhaven National Laboratory using a CCD detector. A single 11bA¹²³⁻³²¹ crystal was used to collect a 2.6Å resolution data set, at 100K, using in-house X-ray on image plate. DATA were processed with DENZO and SCALEPACK to an Rsym of 8.8%, 7.7%, respectively.
30 The starting models were the refined 1.8 Å Mg²⁺ structure (pdb accession code lido) (Lee 1995 *Cell* 80:631) comprising residues D132 to K315, and the refined 2.0 Å Mg²⁺-structure (pdb accession code 1jlm) (Lee et al. 1998 *Structure* 6:923), comprising residues D132 to A318, with the metal and water molecules removed, respectively. The

5 preliminary rigid body refinements were performed using X-plor, with the diffraction data range from 8.0 to 3.0 Å resolution with 5% of reflections for R-free calculation. The phases were gradually extended to high resolution and the structures were refined by several cycles of alternating torsion-angle dynamics and restrained individual isotropic B factor refinement protocols with all diffraction data between 8.0 and 2.6 Å, and 8.0 and 2.3 Å resolution, for 11bA¹²³⁻³²¹ and 11bA¹²³⁻³¹⁵ structures, respectively (Table 1). Model inspection and manual adjustments were made on SGI graphics workstations using O (Jones et al. 1991 *Acta Crystallogr* 47:110). The addition of solvent molecules was based on the suitable peaks in difference maps, reasonable hydrogen bond and refined temperature factors of less than 50 Å². The addition of solvent molecules was based on the suitable peaks in difference maps, reasonable hydrogen bond and refined temperature factors of less than 50 Å². The structure was refined to a final R factor of 20.3% (R-free: 25.6%) for the 11bA¹²³⁻³¹⁵ structure, and 21.9% (R-free: 30.0%) for the 11bA¹²³⁻³²¹ structure. The final models comprises all nonhydrogen atoms of residues D132 to K315, 30 water molecules, and one Ca²⁺ ion for 11bA¹²³⁻³¹⁵, and residues D132 to G321, 44 water molecules and one Mn²⁺ ions for 11bA¹²³⁻³²¹. Crystallographic data is presented in Table 1.

Table 1

		11bA ¹²³⁻³²¹	11bA ¹²³⁻³¹⁵	11bA ^{I316→G}
25	Space group	P2 ₁ 2 ₁ 2	P4 ₃	P4 ₃
	Unit cell constants (Å)	a=48.1, b=121.5, c=74.6	a=b=45.7	
	a=b=45.2, c=95.0		c=94.8	
30	No. of unique reflections	13,990	7,955	3,749
	Resolution (Å)	8.0 - 2.6	8.0 - 2.3	8.0 - 3.0
	Rmerge (%) [*]	7.7 {23} [§]	8.8 {28} [§]	11.2
	{30.4} [§]			
35	Completeness (%)	99.3 {98.4} [§]	91.1 {80.8} [§]	95.3
	{73.6} [§]			
	Redundancy	5.9	2.3	2.7
	R-factor [†] (%)	21.9	18.8	20.8

5	R-free * (%)	30.0	24.8	28.8
	Solvent molecules	82	62	20
	Metal ions	2 (Mn)	1 (Ca)	1 (Ca)
	Root mean square (rms) deviations from ideal values			
	Bond lengths (Å)	0.007	0.007	0.006
10	Bond angle (°)	1.3	1.3	1.25

*R_{merge} = $\sum |I - \langle I \rangle| / \sum I$ where I is the observed intensity and $\langle I \rangle$ is the average intensity from multiple observations of symmetry-related reflections.

§ Numbers in {} shows values for highest 0.1 Å resolution bin.

† R-factor = $\sum |F_o - F_c| / \sum F_o$.

* R-free = $\sum_T |F_o - F_c| / \sum_T F_o$, where T is a test set containing a randomly selected 5% of the reflections omitted from the refinement.

20 The crystal structure 11bA¹²³⁻³²¹ was that of the "closed" conformation (Figures 2A, 2C, and 2E). In contrast, 11bA¹²³⁻³¹⁵ crystallized in the "open" form (Figures 2B, 2D, and 2F).

The ligand binding properties of 11bA¹²³⁻³²¹ and 11bA¹²³⁻³¹⁵ were determined using surface plasmon resonance as previously described (Li et al., *supra*) on a BIAcore (BIAcore AB, Uppsala, Sweden). IC3b, fibrinogen or CD54 each was covalently couples
25 via primary amine groups to the dextran matrix of a separate CM5 sensor chip (BIAcore AB). BSA immobilized in the same way was used as a control surface. 11bA¹²³⁻³²¹, 11bA¹²³⁻³¹⁵ and 11bA^{I-G} A-domains were flowed over the chip at 5 ml/min at different times. TBS (20 mM Tris-HCl, pH 8.0, 150 mM NaCl) with 2 mM MgCl₂ and 0.005%
30 P20 (BIAcore AB) was used as the running buffer. 1 M NaCl in 20 mM Tris-HCl, pH 8.0, was used to remove the bound proteins and to regenerate the surface. Binding was measured as a function of time. The binding data (after subtracting background binding to BSA-coated chip) were analyzed using Scatchard plots as described (Dall'Acqua et al. 1996 *Biochemistry* 35:9667-9676). COS M7 simian fibroblastoid cells at ~70%
35 confluence were transfected with supercoiled cDNAs encoding WT or CD11b^{I-G} together with full-length CD18 as described (Michishita et al., *supra*)

Transfected COS cells were grown for 24 h in Iscove's modified Dulbecco's medium (BioWhittaker, Inc., Walkersville, MD) supplemented with 10% FBS, 2 mM glutamine, 50 IU/ml penicillin and streptomycin at 37°C. Cells were washed, detached

- 5 with 0.1% trypsin-EDTA, and seeded in replicates for 24 h onto 24- or 48-well plates (Costar Corp., Cambridge, MA) or 100-mm petri dishes. Confluent monolayers in 24- or 48-well plates were used for cell-surface antigen quantification and ligand-binding studies, and those on petri dishes for immunoprecipitation studies. Heterodimer formation and binding of iC3b-coated erythrocytes to wild type and CR3^{I→G}
- 10 holoreceptors were carried out as described (Rieu et al. 1996 *J Biol Chem* 271:15858). Specific binding of iC3b to the holoreceptors was obtained by subtracting background binding to mock-transfected COS cells. Binding to CR3^{I→G} was expressed as a percentage of binding to wild type, after correcting for the degree of surface expression using binding of mAb904 (Rieu et al. 1996 *J Biol Chem* 271:15858).
- 15 As shown in Figures 3A-3J, 11bA¹²³⁻³²¹ showed no binding to the activation-dependent "physiologic" ligands, complement iC3b, fibrinogen and CD54 (ICAM-1). In contrast, 11bA¹²³⁻³¹⁵ displayed high affinity binding to all three ligands (Figures 3A-3J). Both domains bound equally well to the activation-independent "ligands" NIF (neutrophil inhibitory factor) (Figures 3D, and 3I), and mAb 904 (Figures 3E and 3J), indicating that
- 20 the differences observed are not caused by variations in A-domain concentrations. These data conclusively establish that the "open" and "closed" crystal structures correspond respectively to the "high" and "low" affinity states of integrin 11bA.

Example 2: Generation of a stable, high affinity state CD11b variant by substitution

- Ile³¹⁶ is invariable in all integrin alpha A-domains cloned to date (Figure 5). An
- 25 A-domain with an Ile to glycine substitution (11bA^{I→G}) exhibited a "high affinity" state (Figure 4A). The same substitution created in the holoreceptor dramatically increased its ligand binding activity (Figure 4B). The crystal structure 11bA^{I→G} was identical to that of the "open" 11bA¹²³⁻³¹⁵ form. These data firmly establish that the "open" high affinity conformation is primarily dictated by an Ile-based switch, intrinsic to the domain, and
- 30 acting allosterically to regulate ligand binding affinity on the MIDAS face. It is known that the inactive and active conformers of an integrin exist in the absence of ligand (Smith et al., *supra*; Yan et al., *J. Biol. Chem.* 275:7249-60, 2000). Based on the results described above, it appears that the role of the ligand is not in initiating the high affinity

5 state as has been recently proposed (Emsely et al, *supra*), but in stabilizing it, through shifting the equilibrium between the low and high affinity state in favor of the latter (Li et al., *supra*). Integrin activation by inside- out signaling would lead to a shift in this equilibrium, increasing the proportion of high affinity receptors on the cell surface that become "available" for ligand binding. Ligand engagement would then generate new
10 epitopes, perhaps extrinsic to the A-domain, that initiate outside-in signaling.

A conserved hydrophobic intramolecular socket (SILEN, Socket for Isoleucine), fastens the I³¹⁶ finger in the "closed" conformation; I³¹⁶ is replaced by L³¹² in the "open" structure (Figures 1A and 1F). SILEN is formed by the hydrophobic side chains of I¹³⁵, L¹⁶⁴, I²³⁶, and Y²⁶⁷ both in the "closed" and "open" conformations. (Figures 1C and 1D).
15 In several integrins, certain mutations that lie outside MIDAS produce gain-of function effects in the holoreceptor, and are believed to act allosterically (Zhang et al., *supra*; Oxvig et al., *supra*; Zhang et al., *supra*). These studies were carried out in the holoreceptors, making it difficult to provide a mechanistic interpretation, because of potential interdomain interactions and/or other quaternary effects. These mutations occur
20 in or around SILEN. For example, substitution of the α 1- β B loop of CD11b with that of CD11a, generates a constitutively active integrin (Zhang et al., *supra*). This region spans one of the SILEN residues L¹⁶⁴. Integrin activation also occurs in an L¹⁶⁴-F substitution (Oxvig et al, *supra*), which predictably makes SILEN smaller and therefore less accommodation to I³¹⁶. Other activating mutations involving E¹³¹, D¹³², K²³¹ and F²³⁴ lie
25 at the bottom of the structure, in close proximity to SILEN (Oxvig et al., *supra*), and may thus exert their effect through interference with the proper coordination of the Ile "finger" in SILEN. The inhibitory effect of certain mAbs with epitopes on the opposite side of MIDAS (e.g., mAb 44a, the epitope of which spans residues on the top of SILEN) may similarly be explained through stabilization of the SILEN pocket.

30 The presence of the N-terminal extension facilitates A-domain switching into the less favored high-affinity state (Li et al, *supra*). The underlying structural basis for this effect is unknown, since none of the residues in the N-terminal extension are included in the derived 3-D structures. It has been observed however that residues within this

5 extension regulate ligand binding in A-domains. First, naturally occurring point mutations in this segment of the vWF A1 domain cause gain-of-function phenotypes in patients with type IIB vWf disease (Matsushita et al., *J. Biol. Chem* 279:13406-14, 1995). This region also contains an activating mutation in CD11b A domain (Oxvig et al., *supra*). Third, synthetic peptide from the N-terminal extension inhibited CD11a-
10 dependent adhesion. Structural data from the CD49b A-domain also show that three residues that extend beyond the $\alpha 7$ helix can pack into a crevice formed in part by residues in the N-terminal extension, bringing the N and C termini into very close proximity (Emsley et al., *supra*). Flexibility of the C-terminal residues in $\alpha 7$ has also been observed in the crystal and NMR structures of the CD11a A-domain. Taken
15 together, these data suggest the N-terminal extension may offer an alternative by imperfect "competitive" surface for luring Ile away from SILEN, allowing some molecules to exist in the "open" form. The present data suggest that such a mechanism may operate in the holoreceptor, providing a mechanistic basis for integrin activation by inside-out signals.

20 Variant Integrin Polypeptides

Given the sequence similarity among the A domains of integrin α subunits, deletion or substitution of the Ile in a selected integrin α subunit that corresponds to Ile³¹⁶ of CD11b should result in the creation of a variant integrin α subunit that is more active (i.e., in solution has a greater proportion of ligand binding form polypeptides) than the
25 wild-type form of the subunit. Figure 5 is an alignment of the C-terminal $\alpha 7$ helix of the A domains of nine integrin α subunits (CD11b, CD11c, CD11d, CD11a, $\alpha 1$, $\alpha 2$, $\alpha 10$, $\alpha 11$, and αE). In this alignment, the invariant Ile corresponding to Ile³¹⁶ of CD11b in the other integrin α subunits is outlined (arrow). Replacing the invariant Ile with Ala or Gly or some other suitable amino acid should create a variant
30 integrin polypeptide with increased activity. We have shown this to be the case in the CD11a A-domain (Figure 6). A variant CD11a A domain containing an I to G substitution displays binding to the activation dependent ligand ICAM-1 in an ELISA

- 5 assay. No binding was observed in the wild-type protein without this substitution. Alternatively, the portion of the integrin α subunit (or the A domain) that includes the invariant Ile and all amino acid residues C terminal to the invariant Ile can be deleted. Table 2 below lists the position of the invariant Ile in each of the integrin α subunits depicted in Figure 5.

10 Table 2

Integrin α Subunit	GenBank Accession No.	Invariant Ile Position in whole integrin	A domain
Human CD11b	RWHU1B	Residue 332	C144-A334
Human CD11c	RWHU1C	Residue 333	C145-A335
Human CD11d	AAB38547	Residue 332	C144-A334
Human CD11a	AAC31672	Residue 331	C150-V333
Human Alpha 1 (CD49a)	P56199	Residue 331	C139-A333
Human Alpha 2 (CD49b)	NP_002194	Residue 361	C169-S363
Human Alpha 10	XP_002097	Residue 249	C57-G251
Human Alpha 11	NP_036343	Residue 349	C159-S351
Human Alpha E	A53213	Residue 385	E196-S387

- The present invention features variant integrin α subunit polypeptides in which the invariant Ile (listed in Table 2) is substituted by Gly, Ala or some other amino acid (e.g., Val). The polypeptide can include part or all of the indicated A domain, e.g., 10, 20, 30, 40, 50, 60, 70, 80, 90, 100, 110, 120, 130, 140, 150, 160, 170, 180, 190, or 200 contiguous amino acids of the A domain that includes the position of the invariant Ile. The invention also includes variant integrin α subunits in which the Ile has been deleted. Also within the invention are polypeptides comprising the entire A domain of an integrin α except for the invariant Ile. For example, amino acids 144-132 of CD11b, amino acids 145-332 of CD11c, amino acids 144-331 of CD11d, amino acids 150-330 of CD11a, but

5 does not include the remainder of the integrin α subunit. The invention also features polypeptides comprising the A domain of an integrin α subunit up to but not including the invariant Ile and further lacking the 5 amino acids following the invariant Ile (e.g., amino acids 144-331, but not 332-336 of CD11b; amino acids 145-332, but not 333-337 of CD11c; amino acids 144-331, but not 332-336 of CD11d; amino acids 150-330, but not 331-335 of CD11a; amino acids 139-330, but not 331-335 of human alpha 1; amino acids 169-360, but not 361-335 of human alpha 2; amino acids 57-248, but not 249-253 of human alpha 10; amino acids 159-348, but not 349-353 of human alpha 11; or amino acids 196-384, but not 385-389 of human alpha E).

15 Given the sequence similarity among the A-like domains of integrin β subunits, deletion or substitution of the Ile in a selected integrin β subunit that corresponds to Ile³¹⁶ of CD11b should result in the creation of a variant integrin β subunit that is more active (i.e., in solution has a greater proportion of ligand binding form polypeptides) than the wild-type form of the subunit. Figure 7 is an alignment of the A-like domains of integrin β subunits. Replacing the conserved Ile with Ala or Gly or some other suitable amino acid should create a variant integrin polypeptide with increased activity. Alternatively, 20 the portion of the integrin β subunit (or the A-like domain of the subunit) that includes the conserved Ile and all amino acid residues C terminal to the invariant Ile can be deleted. Table 3 below lists the position of the conserved Ile in each of the integrin β subunits depicted in Figure 7.

25 Table 3

Integrin β Subunit	GenBank Accession No.	Conserved Ile Position	A-like domain
Human β 2(CD18)	XP_009806	L363	Y125-S365
Human β 1	NP_002202	L378	Y141-S380
Human β 3	I77349	I377	Y136-S379
Human β 4	AAC51632	I366	S128-S368
Human β 5	A38308	I378	Y136-S380

Human β 6	NP_000879	L371	Y131-S373
Human β 7	NP_000880	L389	Y150-S391
Human β 8	NP_002205	L384	Y145-S386

5

Stable open (high affinity) and closed (low affinity) forms of CD11b can also be created by introducing cysteine residues into CD11b so as to create a disulfide bridge that stabilizes the desired form of CD11b (or CD11b A domain). Thus, the invention also features a variant CD11b A domain-containing polypeptide in which F313 and A320 are replaced by cysteine residues (Amino acid positions refer to full-length CD11b starting with a M, not mature CD11b, starting with an F.). This variant forms a stable open form of the A domain. The invention also features a variant CD11b A domain-containing polypeptide in which V315 and A320 are replaced by cysteine residues (Amino acid positions refer to full-length CD11b starting with a M, not mature CD11b, starting with an F.). This variant forms a stable closed form of the A domain. Similar changes can be made in A domain-containing peptides derived from CD11a, CD11c, CD11d, and other integrin subunits. Table 4 indicates the residues that should be changed to cysteines in order to prepare stable closed and open forms of the indicated integrins. The stable open form is useful for generating antibodies directed against active integrins. Active open form polypeptides are also useful for inhibiting the interaction between an integrin and its ligand.

Table 4

Integrin α Subunit	Mutations for Stable Closed Form	Mutations for Stable Open Form
Human CD11b	A320→C V315→C	A320→C F313→C
Human CD11c	A321→C F316→C	A321→C F314→C
Human CD11d	A320→C V315→C	A320→C F313→C
Human CD11a	K319→C L314→C	K319→C K312→C
Human Alpha 1	A319→C	A319→C

(CD49a)	F314→C	F312→C
Human Alpha 2 (CD49b)	A349→C V344→C	A349→C F342→C
Human Alpha 10	A337→C V332→C	A337→C F330→C
Human Alpha 11	A337→C V332→C	A337→C F330→C
Human Alpha E	A373→C V368→C	A373→C V366→C

5

Nucleic Acid Molecules Encoding Variant Integrin Polypeptides

The invention features isolated or purified nucleic acid molecules that encodes a variant integrin polypeptide, e.g., a full length variant integrin subunit or a fragment thereof (e.g., in which the invariant Ile is deleted or substituted), e.g., a biologically active variant integrin polypeptide.

10

In one embodiment, an isolated nucleic acid molecule of the invention includes the nucleotide sequence encoding amino acids 123-315 of the CD11b α subunit (SEQ ID NO:1) or a portion thereof. The nucleic acid molecules can include non-coding sequences or sequences encoding all or a portion of a protein other than an integrin.

15

In another embodiment, an isolated nucleic acid molecule of the invention includes a nucleic acid molecule which is a complement of the nucleotide sequence encoding amino acids 123-315 of the CD11b α subunit.

A nucleic acid molecule of the invention can include only a portion of the nucleic acid sequence of SEQ ID NO:1. For example, such a nucleic acid molecule can include a fragment which encodes all or a portion (e.g., an immunogenic or biologically active portion) or an integrin A domain or A-like domain.

20

Variant Integrin Polypeptides

The present invention also includes variant integrin polypeptides. Such polypeptides can be produced using recombinant DNA methods, by chemical synthesis or using other techniques. The polypeptides can be used as immunogens or antigens

25

5 generate antibodies which bind the active form of an integrin A-domain or A-like domain. The polypeptide can be post-translationally modified, e.g., glycosylated.

A variant integrin polypeptide can be part of a fusion protein which includes all or a portion of a second polypeptide that is not a variant integrin polypeptide. This second polypeptide can be fused to the C-terminus or the N-terminus of the variant integrin polypeptide. All or part of a third peptide may also be present. Thus, a variant integrin polypeptide can be fused to, e.g., GST, an immunoglobulin constant region, a heterologous signal sequence.

The variant integrin polypeptide fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject *in vivo*. For example, the variant integrin polypeptides can be used to reduce skeletal muscle injury. Isolated CD11b A domain has already been found to reduce damage in an animal model of this type of injury. Briefly, purified recombinant rat CD11b A-domain was administered intravenously in a single dose (1mg/kg) to seven groups of Lewis rats (5 per group), 30 minutes before inducing mechanically skeletal muscle injury. Equal numbers of rats were treated with a function-blocking anti-CD11b/CD18 mAb (1 mg/kg). Quantitative histological examination of the wounded area in controlled rats (treated with PBS), showed edema, myofiber disruption, necrosis and erythrocytes extravasation. Influx of neutrophils was detected 30 minutes post wound, followed by a second wave 3 hours later. There was also significant tissue necrosis outside the immediate wounded area (5 mm zone) associated with the presence of activated neutrophils. A-domain or mAb-treated rats showed a comparable and significant decrease in the number of infiltrated PMN ($75 \pm 10\%$, $n=35$) and a protection of the muscular fibers outside the immediate zone of necrosis ($80 \pm 8\%$, $n=35$). These data show that the A-domain can be an effective tissue-preserving agent in this model of muscular injury. Variant A domain should be equally effective.

The variant integrin polypeptide fusion proteins can be used to affect the bioavailability of a variant integrin polypeptide ligand.

5 Variant integrin polypeptide fusion proteins may be useful therapeutically for the treatment of disorders caused by, for example, ischemia-reperfusion injury (*Stroke* 30:134-9,1999), immune complexes (*J Exp Med* 186:1853-63,1997), restenosis, and parasitic diseases (e.g. *Ancylostoma* spp; see *J Cell Biol* 127:2081-91, 1994).

Moreover, the variant integrin polypeptide-fusion proteins of the invention can be
10 used as immunogens to produce anti-variant integrin polypeptide antibodies in a subject, to purify variant integrin polypeptide ligands and in screening assays to identify molecules that inhibit the interaction of variant integrin polypeptide with an variant integrin polypeptide ligand.

Expression vectors are commercially available that already encode a fusion
15 moiety (e.g., a GST polypeptide). An variant integrin polypeptide-encoding nucleic acid can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the variant integrin polypeptide protein.

Variant integrin polypeptides can be produced using an expression vectors (e.g., a plasmid vector or a viral vector). The vector can be autonomously replicating or
20 integrated in to the host genomes. The expression vector can include at least one regulatory sequence (e.g., promoter, enhancer, polyA site, or a cell- or tissue-specific transcription factor binding site) that is operatively linked to the nucleic acid sequence to be expressed. The expression vector can be designed for expression in prokaryotic or eukaryotic cells, e.g., plant cell, insect cells, yeast cells, mammalian cells, and *E coli*.
25 Depending on the host cells used to express the variant integrin polypeptide, it may be desirable to encode the polypeptide using codons optimized for the host cell. In some cases it may be desirable to employ an expression vector capable of directing tissue-specific expression.

The invention also features host cells or recombinant cells harboring a nucleic
30 acid molecule encoding a variant integrin polypeptide. The nucleic acid molecule can be integrated into the host cells genome or present in an autonomously replicating vector. A host cell can be any prokaryotic or eukaryotic cell, e.g., *E. coli*, an insect cell, yeast or

- 5 a mammalian cell. The nucleic acid molecules can be introduced into the host cells through transformation or transfection techniques.

A host cell of the invention can be used to produce (i.e., express) a variant integrin polypeptide by culturing the host cell under conditions such that the polypeptide is produced and then isolating the polypeptide from the cells or the culture medium.

10 Antibodies Recognizing Variant Integrin Polypeptides

- The invention also features antibodies directed against a variant integrin polypeptide. Such antibodies bind to the variant polypeptide with greater affinity than they bind to the corresponding wild-type integrin polypeptide. The antibodies can be generated using standard methods and can be screened by comparing the binding of the
- 15 antibody to the variant integrin polypeptide to binding of the antibody to the corresponding wild-type polypeptide.

- The term "antibody" refers to an immunoglobulin molecule or immunologically active portion thereof, i.e., an antigen-binding portion. Examples of immunologically active portions of immunoglobulin molecules include F(ab) and F(ab')₂ fragments which
- 20 can be generated by treating the antibody with an enzyme such as pepsin. The antibody can be a polyclonal, monoclonal, recombinant, e.g., a chimeric or humanized, fully human, non-human, e.g., murine, or single chain antibody. In a preferred embodiment it has effector function. The antibody can be coupled to a toxin or imaging agent for use in diagnosis of occult inflammation (e.g. in abscess or active atherosclerotic plaques).

- 25 Chimeric, humanized, but most preferably, completely human antibodies are desirable for applications which include repeated administration, e.g., therapeutic treatment (and some diagnostic applications) of human patients.

- The anti-variant integrin polypeptide antibody can be a single chain antibody which can be optionally dimerized or multimerized to generate multivalent antibodies.
- 30 The antibody can be designed to have little or no ability to bind to an Fc receptor.

The antibodies of the invention can be used to detect or purify integrin subunits that are in the active conformation. Thus, they can be used to evaluate the abundance and pattern of expression of an active form of an integrin as part of a clinical testing

- 5 procedure. Detection can be facilitated by coupling (i.e., physically linking) the antibody to a detectable substance (i.e., antibody labeling). Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, and radioactive materials (e.g., horseradish peroxidase, alkaline phosphatase, β -galactosidase, acetylcholinesterase, streptavidin/biotin and avidin/biotin, 10 umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin, luminol, luciferase, luciferin, aequorin, ^{125}I , ^{131}I , ^{35}S and ^3H).

Screening Assays to Identify Compounds that Interact with a Variant Integrin

Polypeptide

- 15 The invention features methods for evaluating a compound for the ability to bind to a variant integrin polypeptide or inhibit the ability of an integrin ligand (e.g., a naturally-occurring integrin ligand) to bind to an variant integrin polypeptide. The methods can include contacting the compound with the variant integrin polypeptide in the presence or absence of a ligand and measuring the ability of the compound or the ligand 20 to bind to the variant polypeptide. The methods can be performed *in vitro*, e.g., in a cell free system, or *in vivo*, e.g., in a two-hybrid interaction trap assay. The method can be used to identify naturally-occurring molecules which interact with an integrin. It can also be used to find natural or synthetic inhibitors of the interaction between an integrin and an integrin ligand.

- 25 The compounds tested can be, e.g., proteins, peptides, peptidomimetics, peptoids, and small molecules. The test compounds can be obtained from combinatorial libraries including: biological libraries; peptoid libraries; spatially addressable parallel solid phase or solution phase libraries; synthetic library methods requiring deconvolution; "one-bead one-compound" libraries.

- 30 Libraries of compounds may be presented in solution (e.g., Houghten 1992 *Biotechniques* 13:412-421), or on beads (Lam 1991 *Nature* 354:82-84), chips (Fodor 1993 *Nature* 364:555-556), bacteria (Ladner U.S. Pat. 5,223,409), spores (Ladner U.S. Pat. 5,223,409), plasmids (Cull et al. 1992 *Proc Natl Acad Sci USA* 89:1865-1869) or on

- 5 phage (Scott and Smith 1990 *Science* 249:386-390); Devlin 1990 *Science* 249:404-406; Cwirla et al. 1990 *Proc. Natl. Acad. Sci.* 87:6378-6382; Felici 1991 *J. Mol. Biol.* 222:301-310; Ladner, *supra.*).

The screening assay can be a cell-based assay in which a cell which expresses an variant integrin polypeptide protein or biologically active portion thereof is contacted
10 with a test compound, and the ability of the test compound to modulate variant integrin polypeptide activity is determined. Determining the ability of the test compound to modulate variant integrin polypeptide activity can be accomplished by monitoring, for example, the ability to bind an integrin ligand.

The ability of the test compound to modulate variant integrin polypeptide binding
15 to a compound, e.g., an integrin ligand or to bind to variant integrin polypeptide can also be evaluated. This can be accomplished, for example, by coupling the compound, e.g., the substrate, with a radioisotope or enzymatic label such that binding of the compound, e.g., the substrate, to variant integrin polypeptide can be determined by detecting the labeled compound, e.g., substrate, in a complex. Alternatively, a variant integrin
20 polypeptide or the integrin ligand can be coupled with a radioisotope or enzymatic label to monitor the ability of a test compound to modulate variant integrin polypeptide binding to an integrin ligand. For example, compounds can be labeled with ^{125}I , ^{35}S , ^{14}C , or ^3H , either directly or indirectly, and the radioisotope detected by direct counting of radioemmission or by scintillation counting. Alternatively, compounds can be
25 enzymatically labeled with, for example, horseradish peroxidase, alkaline phosphatase, or luciferase, and the enzymatic label detected by determination of conversion of an appropriate substrate to product.

In yet another embodiment, a cell-free assay is provided in which an variant integrin polypeptide or biologically active portion thereof is contacted with a test
30 compound and the ability of the test compound to bind to the variant integrin polypeptide or biologically active portion thereof is evaluated. Preferred biologically active portions of the variant integrin polypeptide to be used in assays of the present invention include fragments that bind an integrin ligand.

5 Soluble and/or membrane-bound forms of isolated proteins (e.g., variant integrin polypeptide proteins or biologically active portions thereof) can be used in the cell-free assays of the invention. When membrane-bound forms of the protein are used, it may be desirable to utilize a solubilizing agent. Examples of such solubilizing agents include non-ionic detergents such as n-octylglucoside, n-dodecylglucoside, n-dodecylmaltoside, 10 octanoyl-N-methylglucamide, decanoyl-N-methylglucamide, Triton® X-100, Triton® X-114, Thesit®, Isotridecypoly(ethylene glycol ether)_n, 3-[(3-cholamidopropyl)dimethylamminio]-1-propane sulfonate (CHAPS), 3-[(3-cholamidopropyl)dimethylamminio]-2-hydroxy-1-propane sulfonate (CHAPSO), or N-dodecyl=N,N-dimethyl-3-ammonio-1-propane sulfonate.

15 Cell-free assays involve preparing a reaction mixture of the target gene protein and the test compound under conditions and for a time sufficient to allow the two components to interact and bind, thus forming a complex that can be removed and/or detected.

The interaction between two molecules can also be detected, e.g., using 20 fluorescence energy transfer (FET) (see, for example, Lakowicz et al., U.S. Patent No. 5,631,169; Stavrianopoulos et al., U.S. Patent No. 4,868,103). A fluorophore label on the first, 'donor' molecule is selected such that its emitted fluorescent energy will be absorbed by a fluorescent label on a second, 'acceptor' molecule, which in turn is able to fluoresce due to the absorbed energy. Alternately, the 'donor' protein molecule may 25 simply utilize the natural fluorescent energy of tryptophan residues. Labels are chosen that emit different wavelengths of light, such that the 'acceptor' molecule label may be differentiated from that of the 'donor'. Since the efficiency of energy transfer between the labels is related to the distance separating the molecules, the spatial relationship between the molecules can be assessed. In a situation in which binding occurs between 30 the molecules, the fluorescent emission of the 'acceptor' molecule label in the assay should be maximal. An FET binding event can be conveniently measured through standard fluorometric detection means well known in the art (e.g., using a fluorimeter).

- 5 In another embodiment, determining the ability of the variant integrin polypeptide protein to bind to a target molecule, e.g., an integrin ligand, can be accomplished using real-time Biomolecular Interaction Analysis (BIA) as described above (see, e.g., Sjolander and Urbaniczky (1991) *Anal. Chem.* 63:2338-2345 and Szabo et al. (1995) *Curr. Opin. Struct. Biol.* 5:699-705). "Surface plasmon resonance" or "BIA" detects
- 10 biospecific interactions in real time, without labeling any of the interactants (e.g., BIAcore). Changes in the mass at the binding surface (indicative of a binding event) result in alterations of the refractive index of light near the surface (the optical phenomenon of surface plasmon resonance (SPR)), resulting in a detectable signal which can be used as an indication of real-time reactions between biological molecules.
- 15 In one embodiment, the polypeptide or the test compound is anchored onto a solid phase. The polypeptide/test compound complexes anchored on the solid phase can be detected at the end of the reaction. Preferably, the polypeptide can be anchored onto a solid surface, and the test compound, (which is not anchored), can be labeled, either directly or indirectly, with a detectable labels.
- 20 It may be desirable to immobilize either a variant integrin polypeptide, an anti-variant integrin polypeptide antibody or its target molecule to facilitate separation of complexed from uncomplexed forms of one or both of the proteins, as well as to accommodate automation of the assay. Binding of a test compound to a variant integrin polypeptide protein, or interaction of an variant integrin polypeptide protein with a target
- 25 molecule in the presence and absence of a candidate compound, can be accomplished in any vessel suitable for containing the reactants. Examples of such vessels include microtiter plates, test tubes, and micro-centrifuge tubes. In one embodiment, a fusion protein can be provided which adds a domain that allows one or both of the proteins to be bound to a matrix. For example, glutathione-S-transferase/variant integrin polypeptide
- 30 fusion proteins or glutathione-S-transferase/target fusion proteins can be adsorbed onto glutathione sepharose beads (Sigma Chemical, St. Louis, MO) or glutathione derivatized microtiter plates, which are then combined with the test compound or the test compound and either the non-adsorbed target protein or variant integrin polypeptide protein, and the

5 mixture incubated under conditions conducive to complex formation (e.g., at physiological conditions for salt and pH). Following incubation, the beads or microtiter plate wells are washed to remove any unbound components, the matrix immobilized in the case of beads, complex determined either directly or indirectly, for example, as described above. Alternatively, the complexes can be dissociated from the matrix, and
10 the level of variant integrin polypeptide binding or activity determined using standard techniques.

Other techniques for immobilizing either a variant integrin polypeptide or a target molecule on matrices include using conjugation of biotin and streptavidin. Biotinylated variant integrin polypeptide or target molecules can be prepared from biotin-NHS (N-
15 hydroxy-succinimide) using techniques known in the art (e.g., biotinylation kit, Pierce Chemicals, Rockford, IL), and immobilized in the wells of streptavidin-coated 96 well plates (Pierce Chemical).

In order to conduct the assay, the non-immobilized component is added to the coated surface containing the anchored component. After the reaction is complete,
20 unreacted components are removed (e.g., by washing) under conditions such that any complexes formed will remain immobilized on the solid surface. The detection of complexes anchored on the solid surface can be accomplished in a number of ways. Where the previously non-immobilized component is pre-labeled, the detection of label immobilized on the surface indicates that complexes were formed. Where the previously
25 non-immobilized component is not pre-labeled, an indirect label can be used to detect complexes anchored on the surface; e.g., using a labeled antibody specific for the immobilized component (the antibody, in turn, can be directly labeled or indirectly labeled with, e.g., a labeled anti-Ig antibody).

In one embodiment, this assay is performed utilizing antibodies reactive with
30 variant integrin polypeptide or target molecules but which do not interfere with binding of the variant integrin polypeptide to its target molecule. Such antibodies can be derivatized to the wells of the plate, and unbound target or variant integrin polypeptide protein trapped in the wells by antibody conjugation. Methods for detecting such

5 complexes, in addition to those described above for the GST-immobilized complexes, include immunodetection of complexes using antibodies reactive with the variant integrin polypeptide or target molecule, as well as enzyme-linked assays which rely on detecting an enzymatic activity associated with the variant integrin polypeptide or target molecule.

Alternatively, cell free assays can be conducted in a liquid phase. In such an
10 assay, the reaction products are separated from unreacted components, by any of a number of standard techniques, including chromatography, electrophoresis and immunoprecipitation. In a preferred embodiment, the assay includes contacting the variant integrin polypeptide or biologically active portion thereof with a known compound which binds variant integrin polypeptide to form an assay mixture, contacting
15 the assay mixture with a test compound, and determining the ability of the test compound to interact with an variant integrin polypeptide protein, wherein determining the ability of the test compound to interact with an variant integrin polypeptide includes determining the ability of the test compound to preferentially bind to variant integrin polypeptide or biologically active portion thereof, or to modulate the activity of a target molecule, as
20 compared to the known compound.

To identify compounds that interfere with the interaction between a variant integrin polypeptide and an integrin ligand, a reaction mixture containing the variant integrin polypeptide and the ligand is incubated, under conditions and for a time sufficient, to allow the two products to form complex. In order to test an inhibitory
25 agent, the reaction mixture is provided in the presence and absence of the test compound. The test compound can be initially included in the reaction mixture, or can be added at a time subsequent to the addition of the target gene and its cellular or extracellular binding partner. Control reaction mixtures are incubated without the test compound or with a placebo. The formation of any complexes between the variant integrin polypeptide and
30 integrin ligand is then detected. The formation of a complex in the control reaction, but not in the reaction mixture containing the test compound, indicates that the compound interferes with the interaction. Additionally, complex formation within reaction mixtures containing the test compound and variant integrin polypeptide can also be compared to

5 complex formation within reaction mixtures containing the test compound and the corresponding wild-type integrin polypeptide. These assays can be conducted in a heterogeneous or homogeneous format. Heterogeneous assays involve anchoring either the variant integrin polypeptide or the binding partner (i.e., the integrin ligand) onto a solid phase, and detecting complexes anchored on the solid phase at the end of the
10 reaction. In homogeneous assays, the entire reaction is carried out in a liquid phase. In either approach, the order of addition of reactants can be varied to obtain different information about the compounds being tested. For example, test compounds that interfere with the interaction between the variant integrin polypeptides and the binding partner (e.g., an integrin ligand), e.g., by competition, can be identified by conducting
15 the reaction in the presence of the test substance. Alternatively, test compounds that disrupt preformed complexes, e.g., compounds with higher binding constants that displace one of the components from the complex, can be tested by adding the test compound to the reaction mixture after complexes have been formed. The various formats are briefly described below.

20 In a heterogeneous assay system, either the variant integrin polypeptide or the integrin ligand, is anchored onto a solid surface (e.g., a microtiter plate), while the non-anchored species is labeled, either directly or indirectly. The anchored species can be immobilized by non-covalent or covalent attachments. Alternatively, an immobilized antibody specific for the species to be anchored can be used to anchor the species to the
25 solid surface.

In order to conduct the assay, the partner of the immobilized species is exposed to the coated surface with or without the test compound. After the reaction is complete, unreacted components are removed (e.g., by washing) and any complexes formed will remain immobilized on the solid surface. Where the non-immobilized species is pre-
30 labeled, the detection of label immobilized on the surface indicates that complexes were formed. Where the non-immobilized species is not pre-labeled, an indirect label can be used to detect complexes anchored on the surface; e.g., using a labeled antibody specific for the initially non-immobilized species (the antibody, in turn, can be directly labeled or

5 indirectly labeled with, e.g., a labeled anti-Ig antibody). Depending upon the order of addition of reaction components, test compounds that inhibit complex formation or that disrupt preformed complexes can be detected.

Alternatively, the reaction can be conducted in a liquid phase in the presence or absence of the test compound, the reaction products separated from unreacted
10 components, and complexes detected; e.g., using an immobilized antibody specific for one of the binding components to anchor any complexes formed in solution, and a labeled antibody specific for the other partner to detect anchored complexes. Again, depending upon the order of addition of reactants to the liquid phase, test compounds that inhibit complex or that disrupt preformed complexes can be identified.

15 In an alternate embodiment of the invention, a homogeneous assay can be used. For example, a preformed complex of the variant integrin polypeptide and the integrin ligand is prepared in that either the variant integrin polypeptide or the integrin ligand is labeled, but the signal generated by the label is quenched due to complex formation (see, e.g., U.S. Patent No. 4,109,496 that utilizes this approach for immunoassays). The
20 addition of a test substance that competes with and displaces one of the species from the preformed complex will result in the generation of a signal above background. In this way, test substances that disrupt variant integrin polypeptide-binding partner interaction can be identified.

In yet another aspect, the variant integrin polypeptide proteins can be used as "bait
25 proteins" in a two-hybrid assay or three-hybrid assay (see, e.g., U.S. Patent No. 5,283,317; Zervos et al. (1993) *Cell* 72:223-232; Madura et al. (1993) *J. Biol. Chem.* 268:12046-12054; Bartel et al. (1993) *Biotechniques* 14:920-924; Iwabuchi et al. (1993) *Oncogene* 8:1693-1696; and Brent WO94/10300), to identify other proteins, which bind to or interact with variant integrin polypeptide.

30 A two-hybrid assay can be carried out using a variant A domain as the bait protein. Briefly, the variant A domain is fused to the LexA DNA binding domain and used as bait. The prey is an aptamer library cloned into the active site loop of TrxA expression as a fusion protein with an N-terminal nuclear localization signal, a LexA

5 activation domain, and an epitope tag (Colas et al. 1996 *Nature* 380:548; and Gyuris et al. *Cell* 1993 75:791). Yeast cells are transformed with bait and prey genes. If an aptamer binds to variant A domain, the LexA activation domain is brought into proximity with the LexA DNA binding domain and expression of genes having an appropriately positioned LexA binding site increases.

10 To test this system, yeast strain EGY48 was transformed with the bait plasmid and pSH18-34 (a URA3, 2 μ m plasmid containing the GAL1 promoter fused to lacZ, in which the GAL1 enhancer-like Upstream Activating Sequence (UASG) has been replaced with binding sites for eight LexA dimers). The yeast strain contains two reporter genes, LexAop-LEU2 (replaces the yeast chromosomal LEU2 gene) and
15 LexAop-lacZ, carried on a 2 μ m HIS3⁺ plasmid). Transformed cells were plated on medium containing X-gal. The presence of the bait did not cause the expression of β -galactosidase. The bait by itself also did not activate the LexAop-Leu2 gene in EGY48, since the transformed cells did not grow on Leu deficient plates. Next, nuclear transport of the bait was assessed using the bait plasmid and a pJK101 reporter in a repression type
20 assay. In this assay DNA binding by transcriptionally inert LexA fusion proteins can be detected. The pJK101 reporter is a URA3, 2 μ m plasmid containing most of the UASG. A high affinity binding site for lexA is located between the Gal1 transcription start and the UASG. CD11bA-LexA bait impaired the galactosidase activity of yeast harboring the pJK101 plasmid when grown on galactose medium, indicating that this bait enters the
25 nucleus.

An aptamer library was introduced into the Leu and lacZ reporter-containing yeast cells. Synthesis of library proteins was induced by growing the yeast in galactose medium. In the absence of a suitable prey that binds the bait, the yeast cells does not grow on Leu⁻ medium and have no β -galactosidase activity. A cell expressing a suitable
30 prey will form colonies on Leu⁻ medium and have β -galactosidase activity. Selective galactose (but not glucose) inducible expression allows the Leu and LacZ phenotypes to be unambiguously ascribed to the library protein, diminishing the number of library plasmids that must be excluded by subsequent analysis. Plasmids from positive colonies

5 (i.e., colonies able to grow on galactose, Ura⁻, Trp⁻ His⁻ Leu⁻ plates) were rescued (Hoffman et al. 1987 *Gene* 57:267). Before the respective clones were further characterized, the specificity of their interaction with the bait was tested. This was done by showing that they do not interact with unrelated or nonfunctional baits (e.g., a CD11a A-domain-lexA and 11bA-D242A-lexA respectively), with the DNA-binding domain
10 portion of the bait or nonspecifically with the promoters or other elements of the transcription machinery. The library plasmids were rescued from the galactose-dependent Leu⁺ lacZ⁺ yeast and re-introduced into the original selection strain and into other strains containing different baits. Specific interactors confer the galactose-dependent Leu⁺ and lacZ⁺ phenotype to yeast containing the original bait, but not to
15 yeast containing unrelated baits. An interaction mating assay can be used to test specificity. Briefly, a strain that contains the prey is mated with different yeast strains which express either the original bait protein or the control bait proteins. Reporters should only be active in diploids that contain the original bait (Finley et al. (1994) *Proc. Nat'l Acad. Sci USA* 91:12980).

20 This invention further pertains to novel agents identified by the above-described screening assays. Accordingly, it is within the scope of this invention to further use an agent identified as described herein (e.g., an variant integrin polypeptide modulating agent) in an appropriate animal model to determine the efficacy, toxicity, side effects, or mechanism of action, of treatment with such an agent. Furthermore, novel agents
25 identified by the above-described screening assays can be used for treatments as described herein.

Pharmaceutical Compositions

The variant integrin polypeptides, fragments thereof, as well as anti-variant integrin polypeptide antibodies (also referred to herein as "active compounds") of the
30 invention can be incorporated into pharmaceutical compositions. Such compositions typically include the protein or antibody and a pharmaceutically acceptable carrier. As used herein the language "pharmaceutically acceptable carrier" includes solvents, dispersion media, coatings, antibacterial and anti-fungal agents, isotonic and absorption

5 delaying agents, and the like, compatible with pharmaceutical administration.
Supplementary active compounds can also be incorporated into the compositions.

 A pharmaceutical composition is formulated to be compatible with its intended route of administration. Examples of routes of administration include parenteral, intravenous, intradermal, subcutaneous, oral, inhalation, transdermal, transmucosal, and
10 rectal administration. Solutions or suspensions used for parenteral, intradermal, or subcutaneous application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as
15 ethylenediaminetetraacetic acid; buffers such as acetates, citrates or phosphates and agents for the adjustment of tonicity such as sodium chloride or dextrose. pH can be adjusted with acids or bases, such as hydrochloric acid or sodium hydroxide. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

20 Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. For intravenous administration, suitable carriers include physiological saline, bacteriostatic water, Cremophor EL™ (BASF, Parsippany, NJ) or phosphate buffered saline (PBS). In all
25 cases, the composition must be sterile and should be fluid to the extent that easy syringability exists. It should be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid
30 polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various

5 antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, polyalcohols such as manitol, sorbitol, sodium chloride in the composition. Prolonged absorption of the injectable compositions can be brought about by including in the composition an agent which delays absorption, for
10 example, aluminum monostearate and gelatin.

Sterile injectable solutions can be prepared by incorporating the active compound in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle which contains a
15 basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze-drying which yields a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

20 Oral compositions generally include an inert diluent or an edible carrier. For the purpose of oral therapeutic administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules, e.g., gelatin capsules. Oral compositions can also be prepared using a fluid carrier for use as a mouthwash. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included
25 as part of the composition. The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a
30 sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring.

5 For administration by inhalation, the compounds are delivered in the form of an aerosol spray from pressured container or dispenser which contains a suitable propellant, e.g., a gas such as carbon dioxide, or a nebulizer.

 Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be
10 permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, detergents, bile salts, and fusidic acid derivatives. Transmucosal administration can be accomplished through the use of nasal sprays or suppositories. For transdermal administration, the active compounds are formulated into ointments, salves, gels, or creams as generally known in
15 the art.

 The compounds can also be prepared in the form of suppositories (e.g., with conventional suppository bases such as cocoa butter and other glycerides) or retention enemas for rectal delivery.

 In one embodiment, the active compounds are prepared with carriers that will
20 protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art.
25 The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc. Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) can also be used as pharmaceutically acceptable carriers. These can be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811.

30 It is advantageous to formulate oral or parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subject to be treated; each unit containing a predetermined quantity of active compound calculated to

- 5 produce the desired therapeutic effect in association with the required pharmaceutical carrier.

Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, *e.g.*, for determining the LD50 (the dose lethal to 50% of the population) and the ED50 (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD50/ED50. Compounds which exhibit high therapeutic indices are preferred. While compounds that exhibit toxic side effects may be used, care should be taken to design a delivery system that targets such compounds to the site of affected tissue in order to minimize potential damage to uninfected cells and, thereby, reduce side effects.

The data obtained from the cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED₅₀ with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. For any compound used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. A dose may be formulated in animal models to achieve a circulating plasma concentration range that includes the IC₅₀ (i.e., the concentration of the test compound which achieves a half-maximal inhibition of symptoms) as determined in cell culture. Such information can be used to more accurately determine useful doses in humans. Levels in plasma may be measured, for example, by high performance liquid chromatography.

As defined herein, a therapeutically effective amount of protein or polypeptide (i.e., an effective dosage) ranges from about 0.001 to 30 mg/kg body weight, preferably about 0.01 to 25 mg/kg body weight, more preferably about 0.1 to 20 mg/kg body weight, and even more preferably about 1 to 10 mg/kg, 2 to 9 mg/kg, 3 to 8 mg/kg, 4 to 7 mg/kg, or 5 to 6 mg/kg body weight. The protein or polypeptide can be administered one time per week for between about 1 to 10 weeks, preferably between 2 to 8 weeks, more

5 preferably between about 3 to 7 weeks, and even more preferably for about 4, 5, or 6 weeks. The skilled artisan will appreciate that certain factors may influence the dosage and timing required to effectively treat a subject, including but not limited to the severity of the disease or disorder, previous treatments, the general health and/or age of the subject, and other diseases present. Moreover, treatment of a subject with a
10 therapeutically effective amount of a protein, polypeptide, or antibody can include a single treatment or, preferably, can include a series of treatments.

For antibodies, the preferred dosage is 0.1 mg/kg of body weight (generally 10 mg/kg to 20 mg/kg). If the antibody is to act in the brain, a dosage of 50 mg/kg to 100 mg/kg is usually appropriate. Generally, partially human antibodies and fully human
15 antibodies have a longer half-life within the human body than other antibodies. Accordingly, lower dosages and less frequent administration is often possible. Modifications such as lipidation can be used to stabilize antibodies and to enhance uptake and tissue penetration (e.g., into the brain). A method for lipidation of antibodies is described by Cruikshank et al. ((1997) J. Acquired Immune Deficiency Syndromes and
20 Human Retrovirology 14:193).

The present invention encompasses agents which modulate expression or activity. An agent may, for example, be a small molecule. For example, such small molecules include, but are not limited to, peptides, peptidomimetics (e.g., peptoids), amino acids, amino acid analogs, polynucleotides, polynucleotide analogs, nucleotides, nucleotide
25 analogs, organic or inorganic compounds (i.e., including heteroorganic and organometallic compounds) having a molecular weight less than about 10,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 5,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 1,000 grams per mole, organic or inorganic compounds having a molecular weight
30 less than about 500 grams per mole, and salts, esters, and other pharmaceutically acceptable forms of such compounds.

Exemplary doses include milligram or microgram amounts of the small molecule per kilogram of subject or sample weight (e.g., about 1 microgram per kilogram to about

5 500 milligrams per kilogram, about 100 micrograms per kilogram to about 5 milligrams per kilogram, or about 1 microgram per kilogram to about 50 micrograms per kilogram. It is furthermore understood that appropriate doses of a small molecule depend upon the potency of the small molecule with respect to the expression or activity to be modulated. When one or more of these small molecules is to be administered to an animal (e.g., a
10 human) in order to modulate expression or activity of a polypeptide or nucleic acid of the invention, a physician, veterinarian, or researcher may, for example, prescribe a relatively low dose at first, subsequently increasing the dose until an appropriate response is obtained. In addition, it is understood that the specific dose level for any particular animal subject will depend upon a variety of factors including the activity of the specific
15 compound employed, the age, body weight, general health, gender, and diet of the subject, the time of administration, the route of administration, the rate of excretion, any drug combination, and the degree of expression or activity to be modulated.

An antibody (or fragment thereof) may be conjugated to a therapeutic moiety such as a cytotoxin, a therapeutic agent or a radioactive metal ion. A cytotoxin or
20 cytotoxic agent includes any agent that is detrimental to cells. Examples include taxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, tenoposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs
25 or homologs thereof. Therapeutic agents include, but are not limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BSNU) and lomustine (CCNU), cyclophosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis-dichlorodiamine platinum (II)
30 (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine).

5 The conjugates of the invention can be used for modifying a given biological response, the drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity. Such proteins may include, for example, a toxin such as abrin, ricin A, pseudomonas exotoxin, or diphtheria toxin or a protein such as a
10 cytokine or interleukin.

 Alternatively, an antibody can be conjugated to a second antibody to form an antibody heteroconjugate as described by Segal in U.S. Patent No. 4,676,980.

 The nucleic acid molecules of the invention can be inserted into vectors and used as gene therapy vectors. Gene therapy vectors can be delivered to a subject by, for
15 example, intravenous injection, local administration (see U.S. Patent 5,328,470) or by stereotactic injection (see e.g., Chen et al. (1994) *Proc. Natl. Acad. Sci. USA* 91:3054-3057). The pharmaceutical preparation of the gene therapy vector can include the gene therapy vector in an acceptable diluent, or can comprise a slow release matrix in which the gene delivery vehicle is imbedded. Alternatively, where the complete gene delivery
20 vector can be produced intact from recombinant cells, e.g., retroviral vectors, the pharmaceutical preparation can include one or more cells which produce the gene delivery system.

 The pharmaceutical compositions can be included in a container, pack, or dispenser together with instructions for administration.

25 Animal Models of Processes Involving Integrins

 A model of vascular injury that can be useful in testing potential therapeutic compositions is described by Simon et al. (*J. Clin. Invest.* 2000 105:293). Tang et al. (1997 *J Exp Med* 186:1853) describe CD11b knockout mice which can be useful in various screening assays. An animal model of burn injury may also be useful (*Plast*
30 *Reconstr Surg* 1995 96:1177).

5 Methods of Treatment

The present invention provides for both prophylactic and therapeutic methods of treating a subject at risk of (or susceptible to) a disorder or having a disorder associated with aberrant or unwanted integrin expression or activity.

“Treatment” or “treating a patient”, as used herein, is defined as the application or
10 administration of a therapeutic agent to a patient, or application or administration of a therapeutic agent to an isolated tissue or cell line from a patient, who has a disease, a symptom of disease or a predisposition toward a disease, with the purpose to cure, heal, alleviate, relieve, alter, remedy, ameliorate, palliate, improve or affect the disease, the symptoms of disease or the predisposition toward disease. A therapeutic agent includes,
15 but is not limited to, small molecules, peptides, antibodies, ribozymes and antisense oligonucleotides.

Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD₅₀ (the dose lethal to 50% of the population) and the ED₅₀ (the dose
20 therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD₅₀/ED₅₀. Compounds that exhibit large therapeutic indices are preferred. While compounds that exhibit toxic side effects can be used, care should be taken to design a delivery system that targets such compounds to the site of affected tissue in order to minimize potential
25 damage to uninfected cells and, thereby, reduce side effects.

5

WHAT IS CLAIMED IS:

1 1. An isolated nucleic acid molecule encoding a variant CD11b α subunit having the
2 Ile at position 332 replaced with an amino acid selected from Gly and Ala.

1 2. A variant CD11b α subunit polypeptide having the Ile at position 332 replaced
2 with an amino acid selected from Gly and Ala.

1 3. A isolated nucleic acid molecule encoding a polypeptide consisting of amino acids
2 144 to 331 of CD11b α subunit.

1 4. An isolated nucleic acid molecule encoding a polypeptide comprising amino acids
2 144 to 332 of CD11b α subunit wherein the Ile at amino acid 332 has been replaced by an
3 amino acid selected from Gly and Ala.

1 5. A polypeptide comprising amino acids 144 to 332 of CD11b α subunit wherein the
2 Ile at position 332 has been replaced by an amino acid selected from Gly and Ala.

1 6. A polypeptide comprising amino acids 144 to 331 of CD11b α subunit, the
2 polypeptide not comprising amino acids 332 to 1152 of CD11b α subunit.

1 7. An isolated nucleic acid molecule encoding a variant CD11a α subunit having the
2 Ile at position 331 replaced with an amino acid selected from Gly and Ala.

1 8. A variant CD11a α subunit having the Ile at position 331 replaced with an amino
2 acid selected from Gly and Ala.

1 9. An isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 150 to 330 of CD11a α subunit.

1 10. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 150 to 331 of CD11a α subunit wherein the Ile at amino acid 331 has been replaced by
3 an amino acid selected from Gly and Ala.

1 11. A polypeptide comprising amino acids 150 to 331 of CD11a α subunit wherein
2 the Ile at position 331 has been replaced by an amino acid selected from Gly and Ala.

1 12. A polypeptide comprising amino acids 150 to 330 of CD11a α subunit, the
2 polypeptide not comprising amino acids 331 to 1223 of CD11a.

1 13. A method for determining whether a test compound is a candidate compound for
2 binding to CD11b, comprising:

3 (a) contacting a test compound with a polypeptide comprising amino acids 144 to
4 332 of CD11b α subunit wherein the Ile at amino acid 332 has been replaced by an amino
5 acid selected from Gly and Ala, and

6 (b) determining whether the test compound binds to the polypeptide,
7 wherein a compound which binds to the polypeptide is a candidate compound for
8 binding to CD11b.

1 14. A method for generating an antibody that selectively binds to a polypeptide
2 comprising the open form of CD11b, the method comprising inoculating a non-human
3 mammal with a polypeptide comprising amino acids 144 to 332 of CD11b α subunit wherein
4 the Ile at amino acid 332 has been replaced by an amino acid selected from Gly and Ala; and
5 isolating from the mammal an antibody that selectively binds to a polypeptide
6 comprising the open form of CD11b.

1 15. An antibody that selectively binds to a polypeptide consisting of acids 144 to
2 332 of CD11b α subunit wherein the Ile at amino acid 332 has been replaced by an amino
3 acid selected from Gly and Ala.

1 16. The antibody of claim 15 wherein the antibody is a monoclonal antibody.

1 17. A method for determining whether a test compound is a candidate compound for
2 treating an inflammatory disorder, comprising:

3 (a) contacting a test compound with a polypeptide comprising amino acids 144 to
4 332 of CD11b α subunit wherein the Ile at amino acid 332 has been replaced by an amino
5 acid selected from Gly and Ala, and

6 (b) determining whether the test compound binds to the polypeptide,
7 wherein a compound which binds to the polypeptide is a candidate compound for
8 treating an inflammatory disorder.

1 18. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 320 of CD11b α subunit wherein the Phe at amino acid 313 and the Ala at amino
3 acid 320 have been replaced by Cys.

1 19. A polypeptide comprising amino acids 144 to 320 of CD11b α subunit wherein
2 the Phe at amino acid 313 and the Ala at amino acid 320 have been replaced by Cys.

1 20. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 320 of CD11b α subunit wherein the Val at amino acid 315 and the Ala at amino
3 acid 320 have been replaced by Cys.

1 21. A polypeptide comprising amino acids 144 to 320 of CD11b α subunit wherein
2 the Val at amino acid 315 and the Ala at amino acid 320 have been replaced by Cys.

1 22. A method for determining whether a test compound is a candidate compound for
2 binding to CD11b, comprising:

3 (a) contacting a test compound with a polypeptide comprising amino acids 144 to
4 320 of CD11b α subunit wherein the Phe at amino acid 313 and the Ala at amino acid 320
5 have been replaced by Cys, and

6 (b) determining whether the test compound binds to the polypeptide,
7 wherein a compound which binds to the polypeptide is a candidate compound for
8 binding to CD11b.

1 23. A method for generating an antibody that selectively binds to a polypeptide
2 comprising the open form of CD11b, the method comprising inoculating a non-human
3 mammal with a polypeptide comprising amino acids 144 to 320 of CD11b α subunit wherein
4 the Phe at amino acid 313 and the Ala at amino acid 320 have been replaced by Cys; and
5 isolating from the mammal an antibody that selectively binds to a polypeptide
6 comprising the open form of CD11b.

1 24. An antibody that selectively binds to a polypeptide consisting of amino acids
2 144 to 320 of CD11b α subunit wherein the Phe at amino acid 313 and the Ala at amino acid
3 320 have been replaced by Cys.

1 25. The antibody of claim 24 wherein the antibody is a monoclonal antibody.

1 26. A method for determining whether a test compound is a candidate compound for
2 treating an inflammatory disorder, comprising:

3 (a) contacting a test compound with a polypeptide comprising amino acids 144 to
4 320 of CD11b α subunit wherein the Phe at amino acid 313 and the Ala at amino acid 320
5 have been replaced by Cys, and

6 (b) determining whether the test compound binds to the polypeptide,
7 wherein a compound which binds to the polypeptide is a candidate compound for
8 treating an inflammatory disorder.

1 27. A method for determining whether a test compound is a candidate compound for
2 binding to CD11b, comprising:

3 (a) contacting a test compound with a polypeptide comprising amino acids 144 to
4 320 of CD11b α subunit wherein the Val at amino acid 315 and the Ala at amino acid 320
5 have been replaced by Cys, and

6 (b) determining whether the test compound binds to the polypeptide,
7 wherein a compound which binds to the polypeptide is a candidate compound for
8 binding to CD11b.

1 28. A method for generating an antibody that selectively binds to a polypeptide
2 comprising the open form of CD11b, the method comprising inoculating a non-human
3 mammal with a polypeptide comprising amino acids 144 to 320 of CD11b α subunit wherein
4 the Val at amino acid 315 and the Ala at amino acid 320 have been replaced by Cys; and
5 isolating from the mammal an antibody that selectively binds to a polypeptide
6 comprising the open form of CD11b.

1 29. An antibody that selectively binds to a polypeptide consisting of amino acids
2 144 to 320 of CD11b α subunit wherein the Val at amino acid 315 and the Ala at amino acid
3 320 have been replaced by Cys.

1 30. The antibody of claim 29 wherein the antibody is a monoclonal antibody.

1 31. A method for determining whether a test compound is a candidate compound for
2 treating an inflammatory disorder, comprising:

3 (a) contacting a test compound with a polypeptide comprising amino acids 144 to
4 320 of CD11b α subunit wherein the Val at amino acid 315 and the Ala at amino acid 320
5 have been replaced by Cys, and

6 (b) determining whether the test compound binds to the polypeptide,
7 wherein a compound which binds to the polypeptide is a candidate compound for
8 treating an inflammatory disorder.

1 32. An isolated nucleic acid molecule encoding a variant CD11b α subunit having the
2 Ile at position 332 replaced with an amino acid other than Ile.

1 33. A variant CD11b α subunit polypeptide having the Ile at position 332 replaced
2 with an amino acid other than Ile.

1 34. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 332 of CD11b α subunit wherein the Ile at amino acid 332 has been replaced by
3 an amino acid other than Ile.

- 1 35. A polypeptide comprising amino acids 144 to 332 of CD11b α subunit wherein
2 the Ile at position 332 has been replaced by an amino acid selected from Gly and Ala.
- 1 36. An isolated nucleic acid molecule encoding a variant CD11c α subunit having the
2 Ile at position 333 replaced with an amino acid other than Ile.
- 1 37. A variant CD11c α subunit polypeptide having the Ile at position 333 replaced
2 with an amino acid other than Ile.
- 1 38. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 144 to 332 of CD11c α subunit.
- 1 39. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 333 of CD11c α subunit wherein the Ile at amino acid 332 has been replaced by
3 an amino acid other than Ile.
- 1 40. An isolated nucleic acid molecule encoding a variant CD11d α subunit having the
2 Ile at position 332 replaced with an amino acid other than Ile.
- 1 41. A variant CD11d α subunit polypeptide having the Ile at position 332 replaced
2 with an amino acid other than Ile.
- 1 42. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 144 to 331 of CD11d α subunit.
- 1 43. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 332 of CD11d α subunit wherein the Ile at amino acid 332 has been replaced by
3 an amino acid other than Ile.
- 1 44. An isolated nucleic acid molecule encoding a variant CD49a α subunit having the
2 Ile at position 331 replaced with an amino acid other than Ile.
- 1 45. A variant CD49a α subunit polypeptide having the Ile at position 331 replaced
2 with an amino acid other than Ile.

1 46. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 144 to 330 of CD49a α subunit.

1 47. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 332 of CD49a α subunit wherein the Ile at amino acid 331 has been replaced by
3 an amino acid other than Ile.

1 48. An isolated nucleic acid molecule encoding a variant CD49b α subunit having the
2 Ile at position 361 replaced with an amino acid other than Ile.

1 49. A variant CD49b α subunit polypeptide having the Ile at position 361 replaced
2 with an amino acid other than Ile.

1 50. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 144 to 360 of CD49b α subunit.

1 51. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 144 to 361 of CD49b α subunit wherein the Ile at amino acid 361 has been replaced by
3 an amino acid other than Ile.

1 52. An isolated nucleic acid molecule encoding a variant Alpha 10 α subunit having
2 the Ile at position 249 replaced with an amino acid other than Ile.

1 53. A variant Alpha 10 α subunit polypeptide having the Ile at position 249 replaced
2 with an amino acid other than Ile.

1 54. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 57 to 248 of Alpha 10 α subunit.

1 55. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 57 to 249 of Alpha 10 α subunit wherein the Ile at amino acid 249 has been replacedby
3 an amino acid other than Ile.

1 56. An isolated nucleic acid molecule encoding a variant Alpha 11 α subunit having
2 the Ile at position 349 replaced with an amino acid other than Ile.

1 57. A variant Alpha 11 α subunit polypeptide having the Ile at position 349 replaced
2 with an amino acid other than Ile.

1 58. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 159 to 348 of Alpha 11 α subunit.

1 59. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 159 to 349 of Alpha 11 α subunit wherein the Ile at amino acid 349 has been replaced
3 by an amino acid other than Ile.

1 60. An isolated nucleic acid molecule encoding a variant Alpha E α subunit having
2 the Ile at position 385 replaced with an amino acid other than Ile.

1 61. A variant Alpha 11 α subunit polypeptide having the Ile at position 385 replaced
2 with an amino acid other than Ile.

1 62. A isolated nucleic acid molecule encoding a polypeptide consisting of amino
2 acids 196 to 384 of Alpha 11 α subunit.

1 63. An isolated nucleic acid molecule encoding a polypeptide comprising amino
2 acids 196 to 387 of Alpha 11 α subunit wherein the Ile at amino acid 385 has been replaced
3 by an amino acid other than Ile.

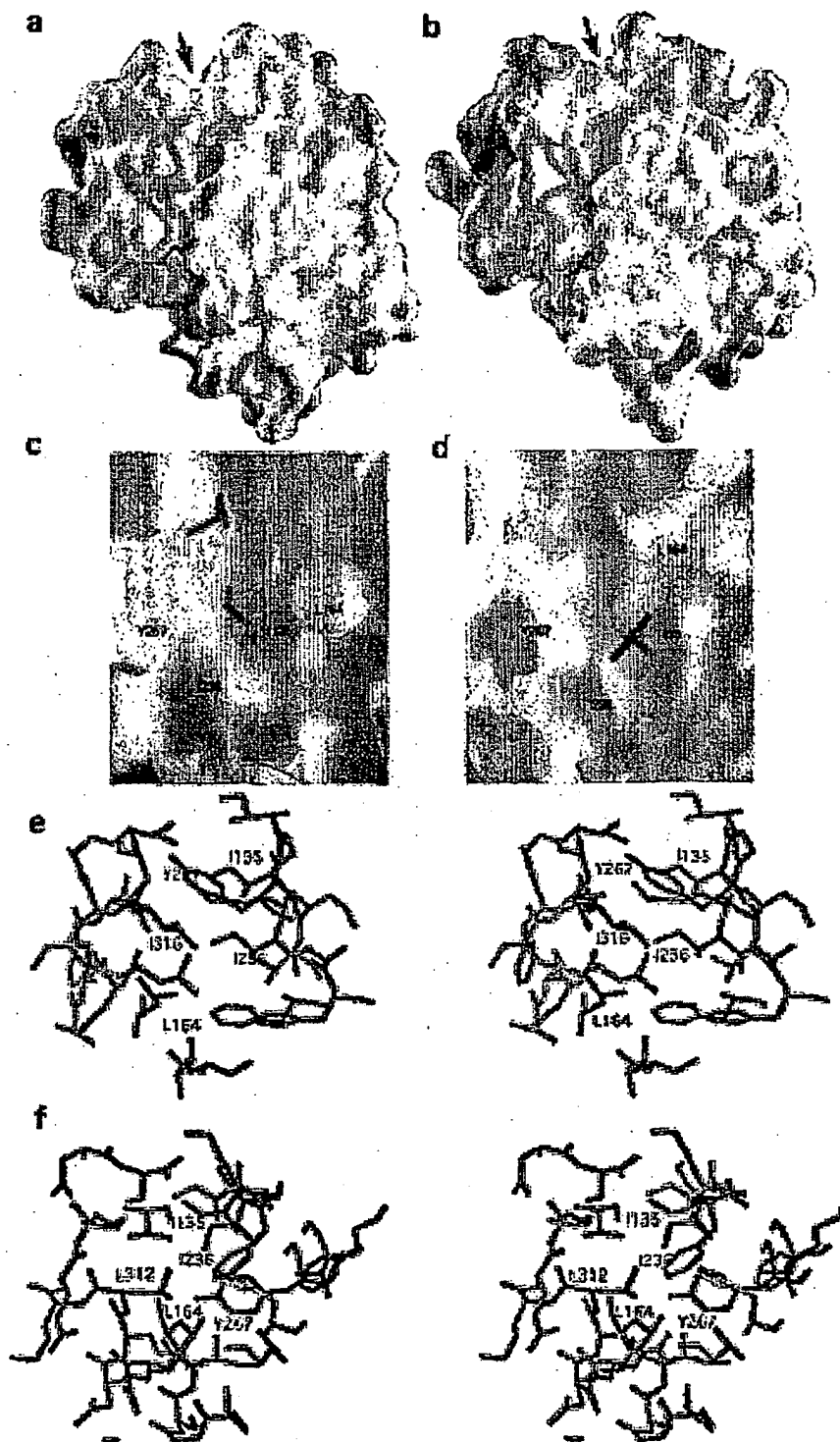
1 64. The isolated nucleic acid molecule of any of claims 32, 34, 36, 38, 39, 40, 42, 43,
2 44, 46, 47, 48, 50, 51, 52, 54, 56, 58, 59, 60, 62, and 63 wherein the amino acids other than
3 Ile is selected from the group consisting of Gly and Ala.

1 65. The polypeptide of any of claims 33, 35, 37, 41, 45, 49, 53, 55, 57, and 61
2 wherein the amino acid other than Ile is selected from the group consisting of Gly and Ala.

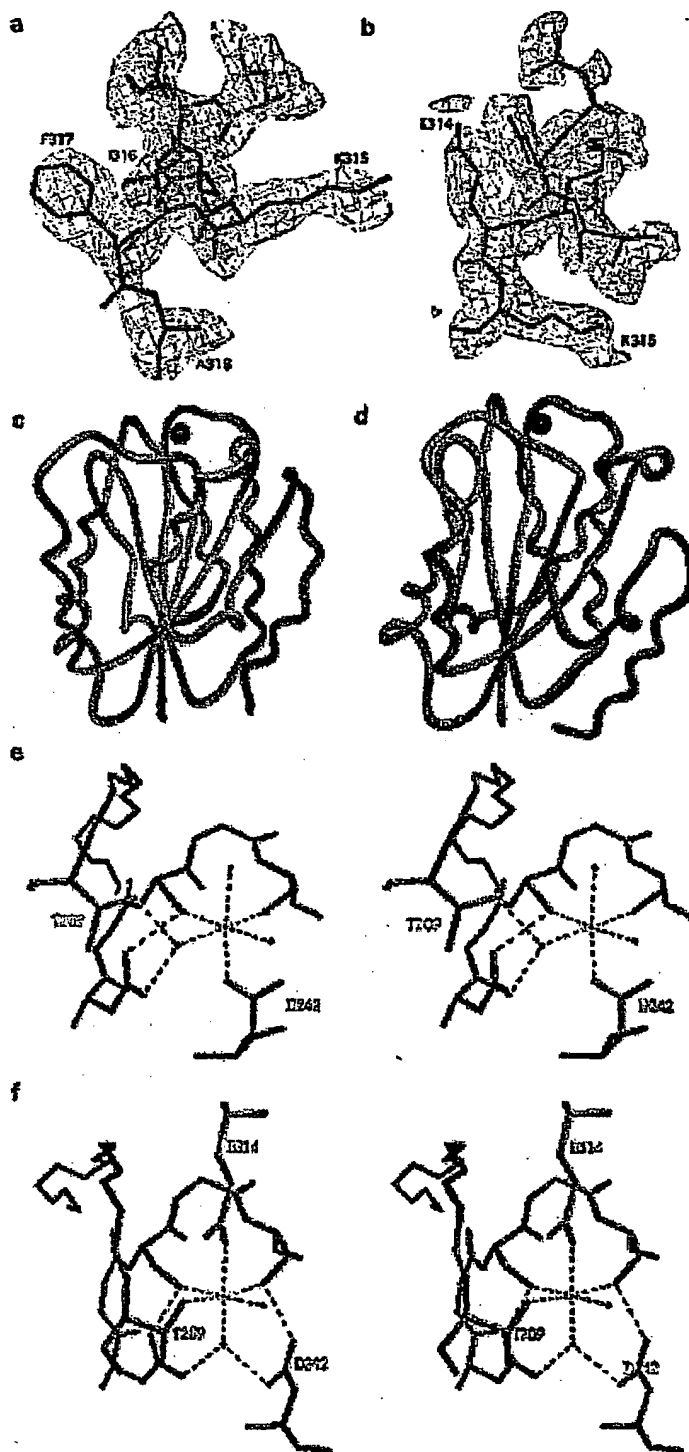
1 66. A method for treating an inflammatory disorder, the method comprising
2 administering the antibody of any of claims 15, 24, and 29.

1 67. A method for treating ischemia-reperfusion injury, the method comprising
2 administering the antibody of any of claims 15, 24, 29.

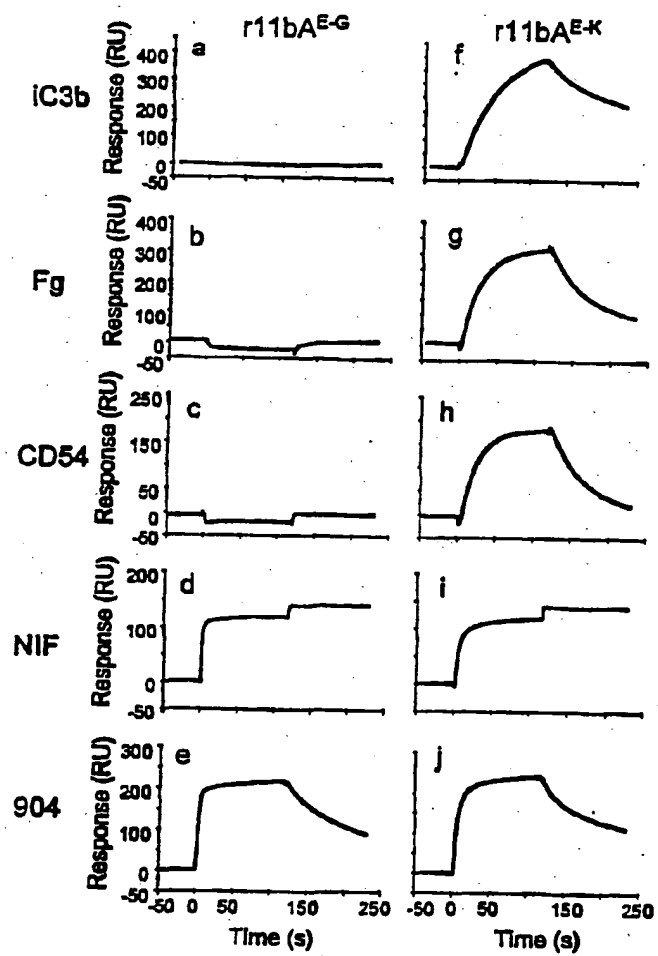
1 68. A method for treating restinosis, the method comprising administering the
2 antibody of any of claims 15, 24, and 29.
3



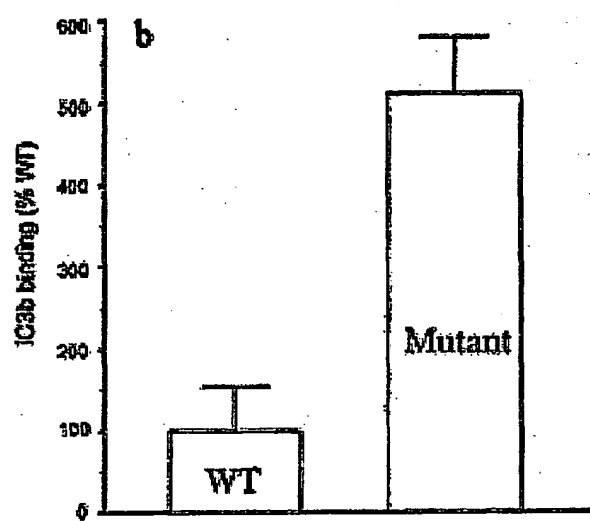
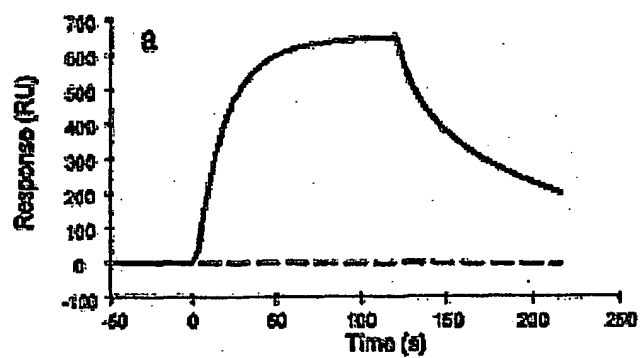
FIGURES 1A-1F



FIGURES 2A-2F



Figures 3A-3J



FIGURES 4A-4B

Docket No.: 00786-536001

[illegible]

FIGURE 5

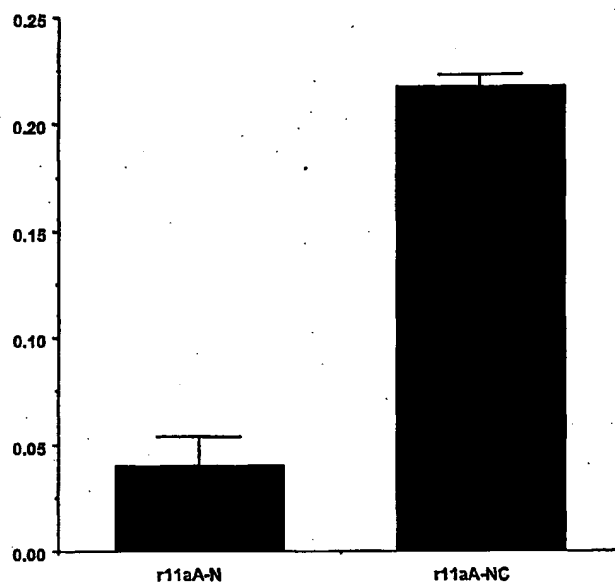


FIGURE 6

b3
b5
b6
b1
b2
b7
b8
b4
con

compendium

	210	220	230	240	250	260	270	280	290	300
b3	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b4	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b5	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b6	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b7	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b8	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b9	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b10	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b11	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b12	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b13	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b14	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b15	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b16	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b17	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b18	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b19	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b20	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b21	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b22	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b23	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b24	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b25	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b26	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b27	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b28	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b29	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b30	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b31	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b32	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b33	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b34	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b35	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b36	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b37	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b38	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b39	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b40	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b41	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b42	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b43	174	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG	ATG
b44	174	ATG	ATG	ATG	ATG</					

[illegible]

→

b3	370	VDA	GR	RA
b5	371	KNA	B	SR
b6	364	167	OE	RR
b1	371	161	GA	S
b2	356	KNA	R	BB
b7	382	TR	A	BB
b8	377	167	GA	CR
b9	359	167	GA	RR
consensus	401	***	***	-

**BY
CONSENT**

REVISED VERSION

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
19 September 2002 (19.09.2002)

PCT

(10) International Publication Number
WO 2002/072030 A2

(51) International Patent Classification⁷: **A61K**

(21) International Application Number:
PCT/US2002/007809

(22) International Filing Date: 13 March 2002 (13.03.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
09/805,354 13 March 2001 (13.03.2001) US

(63) Related by continuation (CON) or continuation-in-part
(CIP) to earlier application:
US 09/805,354 (CIP)
Filed on 13 March 2001 (13.03.2001)

(71) Applicant (for all designated States except US): THE
GENERAL HOSPITAL CORPORATION [US/US]; 55
Fruit Street, Boston, MA 02114 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): ARNAOUT, M.,
Amin [US/US]; 99 Shaw Road, Chestnut Hill, MA 02467
(US). LI, Rui [CN/US]; 89th Street, Apartment 107,
Medford, MA 02155 (US). XIONG, Jian-Ping [CN/US];
95 W. Squantum Street, #1105, Quincy, MA 02171 (US).

(74) Agent: MEIKLEJOHN, Anita, L.; Fish & Richardson
P.C., 225 Franklin Street, Boston, MA 02110 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MY, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SI, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG).

Published:

- with declaration under Article 17(2)(a); without abstract;
title not checked by the International Searching Authority
- with (an) indication(s) in relation to deposited biological
material furnished under Rule 13bis separately from the
description

(48) Date of publication of this revised version:
4 November 2004

(15) Information about Correction:
see PCT Gazette No. 45/2004 of 4 November 2004, Sec-
tion II

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

WO 2002/072030 A2

(54) Title: HIGH AFFINITY INTEGRIN POLYPEPTIDES AND USES THEREOF

(57) Abstract:

PATENT COOPERATION TREATY

PCT

DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT

(PCT Article 17(2)(a), Rule 13ter.1(c) and 39)

Applicant's or agent's file reference G0786-536WO1	IMPORTANT DECLARATION	Date of mailing (day/month/year)
International application No. PCT/US02/07809	International filing date (day/month/year) 13 March 2002 (13.03.2002)	(Earliest) Priority date (day/month/year) 13 March 2001 (13.03.2001)
International Patent Classification (IPC) or both national classification and IPC IPC(7): C07K/ 17/00, 16/00; C07H 21/04; G01N 33/53 and US Cl.: 530/395, 388.1; 536/23.1; 435/7.1		
Applicant THE GENERAL HOSPITAL CORPORATION		

This International Searching Authority hereby declares, according to Article 17(2)(a), that no international search report will be established on the international application for the reasons indicated below.

1. ☒ The subject matter of the international application relates to:
 - a. ☐ scientific theories.
 - b. ☐ mathematical theories.
 - c. ☐ plant varieties.
 - d. ☐ animal varieties.
 - e. ☐ essential biological processes for the production of plants and animals, other than microbiological processes and the products of such processes.
 - f. ☐ schemes, rules or methods of doing business.
 - g. ☐ schemes, rules or methods of performing purely mental acts.
 - h. ☐ schemes, rules or methods of playing games.
 - i. ☐ methods for treatment of the human body by surgery or therapy.
 - j. ☐ methods for treatment of the animal body by surgery or therapy.
 - k. ☐ diagnostic methods practised on the human or animal body.
 - l. ☒ mere presentations of information.
 - m. ☐ computer programs for which this International Searching Authority is not equipped to search prior art.
2. ☒ The failure of the following parts of the international application to comply with prescribed requirements prevents a meaningful search from being carried out:

☐ the description

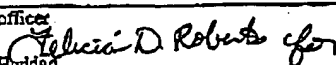
☒ the claims

☐ the drawings
3. ☒ The failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions prevents a meaningful search from being carried out:

☐ the written form has not been furnished or does not comply with the standard.

☒ the computer readable form has not been furnished or does not comply with the standard.

4. Further comments:
Please See Continuation Sheet

Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. 703 305-3230	Authorized officer <div style="text-align: center;">  Maher M. Haddad </div> Telephone No. 703 308-0196
---	---

DECLARATION OF NON-ESTABLISHMENT OF
INTERNATIONAL SEARCH REPORT
Form PCT/ISA/203 (July 1998)

International application No.
PCT/US02/07809

4. Further comments:

Claims 1-68 were unsearchable because the nucleic acid molecules, the polypeptides and the antibodies as well as the variants cannot be matched up in the search. For example, a polypeptide comprising amino acids 150 to 331 of CD11a a subunit wherein the Ile at amino acid 331 has been replaced by amino acid selected from Gly and Ala as in claim 11 is unsearchable because the corresponding amino acids cannot be matched, neither can they be determined from the CD11a a subunit. Also, the claim reads on any CD11a a subunit from any species with any mutation, any substitution, or any deletion. In addition the methods that read on the antibodies and the polypeptides, are unsearchable as well because the active ingredient being used in the claimed methods are unsearchable as noted above.

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☒ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☐ **GRAY SCALE DOCUMENTS**
- ☐ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.